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04/08/2002
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FILE 'REGISTRY' ENTERED AT 15:18:57 ON 08 APR 2002
            348 S (TI AND O)/ELS AND 2/ELC.SUB
T.1
L2
            139 S (TA AND O)/ELS AND 2/ELC.SUB
              5 S (W AND S AND N)/ELS AND 3/ELC.SUB
L3
              4 S (TA AND AL AND O AND N)/ELS AND 4/ELC.SUB
T.4
             43 S (TA AND AL AND O)/ELS AND 3/ELC.SUB
L5
            141 S (AL AND O AND N)/ELS AND 3/ELC.SUB
L6
              1 S PT/CN
L7
L8
              3 S AU/CN
              2 S MO/CN
L9
              3 S TA/CN
L10
              1 S IRIDIUM/CN
L11
L12
              1 S RU/CN
              3 S CR/CN
   FILE 'HCAPLUS' ENTERED AT 15:22:19 ON 08 APR 2002
        439968 S PT OR PLATINUM OR AU OR GOLD
         585582 S MOLYBDENUM OR MO OR TANTALUM OR TA
L15
         533308 S IRIDIUM OR IR OR RUTHERNIUM OR RU
L17
         459264 S CHROMIUM OR CR
          32613 S EMITTER OR ECL
L18
           4571 S (CATHODE (2N) LAYER) OR (FUSED(2N) ELECTROLYTE)
L19
             60 S ELETRON
L20
          4558 S (TUNNEL?) (3N) (FILM? OR LAYER? OR COAT####)
L21
         225148 S (METAL#### OR ALLOY? OR AMALGAM? OR INGOT? OR BULLION?) (5N) (D
L22
           123 S L1 AND L18
L23
             36 S L2 AND L18
L24
              0 S L3 AND L18
L25
              0 S L4 AND L18
L26
             0 S L5 AND L18
L27
             1 S L6 AND L18
L28
             25 S L23 AND L22
L29
L30
             1 S L23 AND L21
L31
             0 S L30 NOT L29
L32
            144 S L18 AND L19
L33
              1 S L32 AND L21
L34
              0 S L32 AND L20
          75276 S TITANIUM(2N)MONOXIDE OR TITANIUM(2N)OXIDE
L35
L36
            109 S L35 AND L18
              1 S TUNGSTEN NITRIDE SULFIDE
L37
              4 S ALUMINUM TANTALUM NITRIDE OXIDE
L38
           119 S ALUMINUM TANTALUM OXIDE
L39
           1160 S ALUMINUM NITRIDE OXIDE
L40
L41
              0 S TANTALAM OXIDE
              0 S L37 AND L18
L42
          13414 S TANTALUM OXIDE
L43
              0 S L38 AND L18
L44
L45
              0 S L39 AND L18
              0 S L40 AND L18
L46
L47
             52 S L43 AND L18
             24 S L36 AND (L21 OR L22)
L48
             11 S L48 NOT L29
L49
L50
           5947 S L18 AND (L7-17)
L51
             38 S L50 AND L19
L52
             37 S L51 NOT (L49 OR L29)
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Serial No.:09/846,127

▶04/08/2002

L30 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2002 ACS

1993:615789 HCAPLUS AN

DN 119:215789

Insulating channel dielectric-base transistor TI

Tamura, Yasutaka IN

PΑ Fujitsu Ltd, Japan

Jpn. Kokai Tokkyo Koho, 8 pp. so

CODEN: JKXXAF

DTPatent

Japanese LA

FAN.CNT 1

APPLICATION NO. DATE PATENT NO. KIND DATE -----

JP 05198854 A2 19930806 JP 1992-9026 19920122 PΙ

The transistor comprises a base region having a high dielec. const., an ΑB emitter electrode on 1 side of the base region through a 1st barrier layer having a dielec. const. lower than that of the base region, an collector electrode through a 2nd barrier layer of the same type, and a base electrode on the other side through a 3rd barrier layer with height and thickness to prevent carrier tunneling. The 3rd barrier layer may have the dielec. const. higher than those of the 1st and 2nd barrier layers. The transistor may comprise an insulating substrate successively covered with the dielec. base region layer, the 3rd barrier layer and the base electrode. The base electrode may be made of an impurity-doped or quasi-stoichiometric base region material. The emitter, collector, and/or base electrode may consist of metal or oxide superconductor. The base region may consist of oxide contg. Sr, Ti, K, Ta, Sn, Zr, or Nb. The base region may consist of KTal-xNbxO3, having controlled temp. of the max. dielec. const. with the Nb content.

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L29 ANSWER 1 OF 25 HCAPLUS COPYRIGHT 2002 ACS
     2002:158258 HCAPLUS
AN
     136:209298
DN
     Process for producing electron emission cathode
ΤI
    Miyamoto, Hiroyuki; Iseki, Misao; Arai, Manabu; Tamai, Hideaki; Kimura,
IN
    New Japan Radio Co., Ltd., Japan
PΑ
    U.S. Pat. Appl. Publ., 20 pp.
SO
    CODEN: USXXCO
DT
    Patent
    English
LA
FAN.CNT 1
                    KIND DATE
                                         APPLICATION NO. DATE
     PATENT NO.
     _____
    US 2002024280
                                         US 2001-934212 20010821
                    A1 20020228
     JP 2002075165
                    A2 20020315
                                         JP 2000-262091 20000831
    DE 10142396
                    A1 20020314
                                         DE 2001-10142396 20010830
    US 2002024281
                    A1 20020228
                                         US 2001-971226 20011003
PRAI JP 2000-262091
                   Α
                           20000831
    US 2001-934212
                    A2
                           20010821
    There is provided a cathode which is easily operable, harmless, and stable
     at high temp. .gtoreq.1,400.degree. as well as excellent in electron
     emission characteristics at the same time, and the process for prepg. the
           The cathode of the present invention comprises a polycryst.
     substance or a polycryst. porous substance of high-m.p. metal material and
     an emitter material dispersed into the polycryst. substance or
    polycryst. porous substance, in which 0.1-30% by wt. of .gtoreg.1 selected
     from the group consisting of Hf oxide, Zr oxide, La oxide, Ce oxide and Ti
    oxide is dispersed in the emitter material.
    Molding
IT
        (isostatic pressing, cold; process for producing electron emission
       cathode)
    Electron emission
ΙT
        (process for producing electron emission cathode)
     11126-28-6, Titanium tungstate 12737-23-4, Cerium tungsten oxide
IT
     37382-36-8, Lanthanum tungstate
                                     39290-95-4, Zirconium tungstate
     51680-39-8, Hafnium tungsten oxide
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PYP (Physical process); TEM (Technical or engineered material
    use); PROC (Process); USES (Uses)
        (compd. layer on electron emission surface; process for producing
       electron emission cathode)
IT
     1333-74-0, Hydrogen, processes
     RL: CPS (Chemical process); NUU (Other use, unclassified); PEP (Physical,
     engineering or chemical process); PROC (Process); USES (Uses)
        (dispersion of emitter materials in atm. of; process for
       producing electron emission cathode)
IT
     1314-35-8, Tungsten oxide, processes
                                           7439-98-7, Molybdenum, processes
     7440-33-7, Tungsten, processes
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PYP (Physical process); TEM (Technical or engineered material
    use); PROC (Process); USES (Uses)
        (emitter materials; process for producing electron emission
       cathode)
IT
    11130-73-7, Tungsten carbide 12627-57-5, Molybdenum carbide
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PYP (Physical process); TEM (Technical or engineered material
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use); PROC (Process); USES (Uses)

04/08/2002 (layer on electron emission surface; process for producing electron emission cathode) 1312-81-8, Lanthanum oxide 1314-23-4, Zirconium oxide, processes IT 7439-88-5, Iridium, processes 7440-04-2, Osmium, processes Rhenium, processes 7440-18-8, Ruthenium, processes 11129-18-3, Cerium 12055-23-1, Hafnium oxide 13463-67-7, Titanium oxide (TiO2), processes RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES (Uses) (mixed with emitter materials; process for producing electron emission cathode) L29 ANSWER 2 OF 25 HCAPLUS COPYRIGHT 2002 ACS ΑN 2001:792159 HCAPLUS 135:337040 DN Liquid crystal display having swithing device driven by ultraviolet beam TIirradiation Kido, Masami INPASharp Corp., Japan Jpn. Kokai Tokkyo Koho, 10 pp. SO CODEN: JKXXAF DT Patent LΑ Japanese FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ______ -----JP 2001305572 A2 20011031 JP 2000-119375 20000420 PΙ AB The display device has an elec. switching device involving a metal oxide layer with energy band gap in UV region, i.e., .gtoreq.3.0 eV, and the metal oxide layer is irradiated with UV from a stripe light source in the device so that the elec. current is regulated for switching elec. voltage onto the liq. crystal. Alternatively, the metal oxide layer is assocd. with another metal oxide layer as a barrier layer, on which a diode involving an anode and a cathode is formed for regulation of elec. current. A transistor may be formed with an emitter electrode and a collector electrode on the barrier metal oxide layer and a base electrode on the former metal oxide layer, wherein elec. current between the emitter and the collector is regulated by elec. voltage on the base electrode under irradn. on the former metal oxide. The voltage on the liq. crystal may be regulated by metal-insulator-metal (MIM) structure on the former metal oxide layer. ITElectric switches Liquid crystal displays UV radiation (display device having UV irradn.-driven switching device for applying elec. voltage on liq. crystal) TΤ Diodes Thin film transistors Transistors (in display device having UV irradn.-driven switching device for

applying elec. voltage on liq. crystal) IT Electroluminescent devices

(light source; in display device having UV irradn.-driven switching device for applying elec. voltage on liq. crystal)

7631-86-9, Silica, uses 12060-59-2, Strontium titanium oxide (SrTiO3) IT 13463-67-7, Titania, uses

```
RL: DEV (Device component use); USES (Uses)
        (film; in display device having UV irradn.-driven switching device for
       applying elec. voltage on liq. crystal)
    ANSWER 3 OF 25 HCAPLUS COPYRIGHT 2002 ACS
L29
AN
    2000:667007 HCAPLUS
    133:225600
DN
    An aluminum alloy back junction solar cell and a process for fabrication
TI
IN
    Meier, Daniel L.; Davis, Hubert P.; Garcia, Ruth A.; Salami, Jalal
    Ebara Solar, Inc., USA
PΑ
     PCT Int. Appl., 39 pp.
SO
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                         APPLICATION NO. DATE
     -----
                                         -----
    WO 2000055923
                    A1 20000921
                                        WO 2000-US2609 20000201
PΙ
        W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
            DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
            JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
            MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
            TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD,
            RU, TJ, TM
        RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
            DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
            CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                     B1 20010717
    US 6262359
                                        US 1999-414990 19991007
    BR 2000009085
                           20020102
                      Α
                                         BR 2000-9085
                                                          20000201
    EP 1166367
                      A1
                           20020102
                                         EP 2000-930073
                                                          20000201
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO
    TW 449933 .
                     В
                           20010811
                                         TW 2000-89102645 20000216
PRAI US 1999-124797P
                      Ρ
                           19990317
    US 1999-414990
                      Α
                           19991007
    WO 2000-US2609
                           20000201
                      W
     A process for fabricating a solar cell includes: providing a base layer,
AB
     and fabricating an emitter layer of p-type cond. on a same side
     as the non-illuminated surface of the base layer to provide a strongly
     doped p-type emitter layer and a p-n junction between the n-type
     base layer and the p-type emitter layer. The base layer of the
     present invention has n-type cond. and is defined by an illuminated
     surface and a non-illuminated surface which is opposite to the illuminated
     surface.
    Bridgman crystal growth
ΙT
     Czochralski crystal growth
     Electric contacts
     Screen printing
     Solar cells
        (aluminum alloy back junction solar cell and process for fabrication
       thereof)
ΙT
     Casting of metals
        (directional solidification; aluminum alloy back junction solar cell
       and process for fabrication thereof)
IT
     Crystal growth
        (edge-defined film-fed; aluminum alloy back
        junction solar cell and process for fabrication thereof)
IT
     Crystal growth
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(floating-zone; aluminum alloy back junction solar cell and process for
        fabrication thereof)
     7440-21-3, Silicon, uses
                               7440-22-4, Silver, uses
                                                         11145-27-0
ΙT
     RL: DEV (Device component use); USES (Uses)
        (aluminum alloy back junction solar cell and process for fabrication
        thereof)
IT
     7429-90-5, Aluminum, uses
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (aluminum alloy back junction solar cell and process for fabrication
IT
     7723-14-0, Phosphorus, uses
     RL: MOA (Modifier or additive use); USES (Uses)
        (aluminum alloy back junction solar cell and process for fabrication
IT
     12033-89-5, Silicon nitride, uses 13463-67-7, Titania, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (antireflective coating; aluminum alloy back junction solar cell and
       process for fabrication thereof)
    11099-22-2
IT
     RL: DEV (Device component use); USES (Uses)
        (eutectic; aluminum alloy back junction solar cell and process for
       fabrication thereof)
             THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 4
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 4 OF 25 HCAPLUS COPYRIGHT 2002 ACS
    2000:191404 HCAPLUS
AN
DN
    132:243723
    Cathodoluminescent solid-state lasers
TT
    Leksono, Moeljanto W.; Qiu, Chang-hua; Pankove, Jacques Isaac
ΙN
PA
    Astralux, Inc., USA
SO
    PCT Int. Appl., 32 pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 2
    PATENT NO.
                    KIND DATE
                                          APPLICATION NO. DATE
     ---- ---- ----
                           _____
                                          -----
PΤ
    WO 2000016454
                     A1
                           20000323
                                          WO 1999-US21585 19990917
        W: CA, JP, MX
        RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
            PT, SE
    US 6067308
                           20000523
                                          US 1998-154813
                      Α
                                                           19980917
PRAI US 1998-154813
                           19980917
                      Α
    Solid-state lasers are described which comprise an elongated, single
     crystal metal oxide body selected from the group Al203
     (sapphire), ZnO, MgO, LiNbO3, TiO3, SrTiO3, BaTiO3, and quartz doped with
     .gtoreq.1 of erbium, terbium, praseodymium, neodymium, samarium, europium,
     dysprosium, holmium, thulium, and ytterbium and with a rare earth-ionizing
    element selected from the group oxygen and fluorine (which ionize the
     .gtoreq.1 rare earth elements); an elongated electron emitter
```

The means to render the emitted radiation coherent may be a Fabry-Perot cavity that includes the metal oxide body or an acoustic generator that is assocd. with the metal oxide body in a manner to produce a standing wave within the metal oxide body. Solid state lasers IT (cathodoluminescent solid-state lasers) Rare earth metals, uses IT RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses) (cathodoluminescent solid-state lasers) IT Visible lasers (solid-state; cathodoluminescent solid-state lasers) 1309-48-4, Magnesium oxide (MgO), uses 1314-13-2, Zinc oxide (ZnO), uses TT 1344-55-4, Titanium trioxide 7631-86-9, Silica, uses 12031-63-9, Lithium niobate 12047-27-7, Barium titanate, uses 12060-59-2, Strontium titanate 13463-67-7, Titania, uses RL: DEV (Device component use); USES (Uses) (cathodoluminescent solid-state lasers) 7429-91-6, Dysprosium, uses 7440-00-8, Neodymium, uses 7440-10-0, Praseodymium, uses 7440-19-9, Samarium, uses 7440-27-9, Terbium, uses 7440-30-4, Thulium, uses 7440-52-0, Erbium, uses 7440-53-1, Europium, 7440-64-4, Ytterbium, uses 7782-41-4, 7440-60-0, Holmium, uses 7782-44-7, Oxygen, uses Fluorine, uses RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses) (cathodoluminescent solid-state lasers) 18195-92-1, Erbium +2, uses 18472-30-5, Erbium +3, uses IT RL: DEV (Device component use); MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (cathodoluminescent solid-state lasers) IT 1344-28-1, Alumina, uses RL: DEV (Device component use); USES (Uses) (sapphire-type; cathodoluminescent solid-state lasers) THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT ALL CITATIONS AVAILABLE IN THE RE FORMAT ANSWER 5 OF 25 HCAPLUS COPYRIGHT 2002 ACS L29 AN 1998:790410 HCAPLUS 130:60002 DN Multilayer composite electrodes for discharge lamps TIMehrotra, Vivek; Betrabet, Hemant S.; McGee, Susan; McGee, Thomas F. INPhilips Electronics North America Corporation, USA PΑ SO U.S., 13 pp. CODEN: USXXAM Patent English

DTLA

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE --------------US 1994-363184 PΙ US 5847498 Α 19981208 19941223 US 1997-832895 19970403 US 5847497 Α 19981208 PRAI US 1994-363184 19941223

Composite sintered electrodes with improved properties that make them suitable for use in a variety of lamp types are provided which comprise a refractory metal and a substantial amt. of a refractory emitter oxide, either single layer or multilayer, the composites having been subjected to sintering at an elevated temp. effective to form a composite electrode having a d. of .gtoreq.85%, preferably in the presence of a

Serial No.:09/846,127

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04/08/2002
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sintering activator, e.g., Ni, or its mixt. with a sintering aid, e.g.,
    Li20.
    Sintering aids
IT
        (for multilayer composite electrodes for discharge lamps)
TΤ
    Electric discharge lamps
    Electrodes
        (multilayer composite electrodes for discharge lamps)
    Alkali metal oxides
IT
    Group VIII elements
    Refractory metal oxides
    Refractory metals
    RL: DEV (Device component use); USES (Uses)
        (multilayer composite electrodes for discharge lamps contg.)
IT
    Sintering
        (of multilayer composite electrodes for discharge lamps)
    7440-33-7, Tungsten, uses 12009-21-1, Barium zirconate
ТТ
                                                               12009-63-1,
    Barium titanium oxide (Ba2TiO4) 12047-27-7, Barium titanate, uses
    13463-67-7, Titanium dioxide, uses 66103-41-1, Barium tantalum
           120898-04-6, Barium strontium zirconium oxide (Ba0.5Sr0.5ZrO3)
    132826-58-5, Barium yttrium oxide
    RL: DEV (Device component use); USES (Uses)
        (multilayer composite electrodes for discharge lamps contg.)
IT
    7440-02-0, Nickel, processes
    RL: PEP (Physical, engineering or chemical process); PROC (Process)
        (sintering activator; for multilayer composite electrodes for discharge
       lamps)
    12057-24-8, Lithium oxide (Li20), processes
IT
    RL: PEP (Physical, engineering or chemical process); PROC (Process)
        (sintering aid; for multilayer composite electrodes for discharge
       lamps)
      19
             THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L29
    ANSWER 6 OF 25 HCAPLUS COPYRIGHT 2002 ACS
    1998:545669 HCAPLUS
AN
DN
    129:182053
    Silver halide photosensitive material and sensor for monitoring its
TI
    movement in automatic conveyer
    Ito, Hirohide; Fukui, Makoto
IN
    Konica Co., Japan
PA
    Jpn. Kokai Tokkyo Koho, 15 pp.
SO
    CODEN: JKXXAF
DT
    Patent
LΑ
    Japanese
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                         APPLICATION NO. DATE
    -----
                                          -----
                    A2 19980821
PΙ
    JP 10221809
                                          JP 1997-23802
                                                           19970206
    The material comprises a transparent support having thereon a
AΒ
    light-sensitive Ag halide emulsion layer with Ag content <4.0 g/m2 and a
    layer contg. particles of .gtoreq.1 metallic oxide
    selected from In203, SnO2, ZnO, Al2O3, and TiO2 having n .gtoreq.1.50 and
    0.2-3.0 .mu.m diam., which scatters IR rays and transmits a visible light.
    The monitoring method is characterized by the followings: (a) an IR sensor
    comprising a pair of an IR ray emitter and an IR detector is
    used; (B) IR ray from the emitter is scattered by the material;
     (C) increase of scattered ray is monitored by the detector. A photog.
    material with low Ag content can be detected by the method without
    affecting the characteristics of the material.
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ΙT IR detectors (for monitoring movement of photog. film having emulsion-protecting layer contg. metal oxide for IR scattering in automatic conveyer) IT Photographic films (photog. film having emulsion-protecting layer contg. metal oxide for IR scattering) Transparent materials IT (support; photog. film having emulsion-protecting layer contg. metal oxide for IR scattering) 1314-13-2, Zinc oxide, uses 1312-43-2, Indium oxide (In2O3) IT Alumina, uses 13463-67-7, Titania, uses 18282-10-5, Tin oxide (SnO2) RL: TEM (Technical or engineered material use); USES (Uses) (particle; photog. film having emulsion-protecting layer contg. metal oxide for IR scattering) ANSWER 7 OF 25 HCAPLUS COPYRIGHT 2002 ACS L29 1998:421628 HCAPLUS ANDN129:126317 Super cleaning of closed space by UV/photoelectron method using TIphotocatalyst Fujii, Toshiaki; Suzuki, Tsukuru; Sakamoto, Kazuhiko; Yokoyama, Shin; ΑU Hirose, Masataka Ebara Res. Co., Ltd., Fujisawa, 251-8502, Japan CS Earozoru Kenkyu (1998), 13(2), 110-118 SO CODEN: EAKEEA; ISSN: 0912-2834 PBNippon Earozoru Gakkai DTJournal LA Japanese A new super cleaning equipment of a closed space using a photocatalyst AΒ (TiO2) in a UV/photoelectron method was developed. The newly developed equipment has photocatalyst between photoelectron emitter and electrode (charging space) in the UV/photoelectron equipment to remove gaseous contaminants, such as hydrocarbons. The equipment developed in this work is characterized as follows:. 1) It can create a super clean space in which both gaseous contaminants, such as hydrocarbons, ammonia and particles are removed simultaneously, 2) when Si wafer or metallic substrates are set in the clean space inside such an equipment, surface contamination of the substrates is prevented, 3) in evaluation of actual electron device using MOS (Metal-Oxide-Semiconductor) capacitor, the reliability of gate oxides is improved in the time dependent dielec. breakdown (TDDB) characteristics. IT Semiconductor materials (manuf. of; super cleaning of closed space by UV/photoelectron method using photocatalyst) IT Air purification (photocatalytic; super cleaning of closed space by UV/photoelectron method using photocatalyst) IT Airborne particles (super cleaning of closed space by UV/photoelectron method using photocatalyst) IT Hydrocarbons, processes RL: REM (Removal or disposal); PROC (Process) (super cleaning of closed space by UV/photoelectron method using photocatalyst) 13463-67-7, Titanium oxide (TiO2), uses IT

(super cleaning of closed space by UV/photoelectron method using

RL: CAT (Catalyst use); USES (Uses)

SO

Eur. Pat. Appl., 47 pp.

photocatalyst) 7664-41-7, Ammonia, processes RL: REM (Removal or disposal); PROC (Process) (super cleaning of closed space by UV/photoelectron method using photocatalyst) ANSWER 8 OF 25 HCAPLUS COPYRIGHT 2002 ACS L29 1997:571375 HCAPLUS AN 127:170256 DN Electron emitters, electron sources, and display devices TIShibata, Masaaki; Tsukamoto, Takeo; Iwasaki, Tatsuya TN PΑ Canon K. K., Japan SO Jpn. Kokai Tokkyo Koho, 23 pp. CODEN: JKXXAF DT Patent LΑ Japanese FAN.CNT 1 KIND DATE PATENT NO. APPLICATION NO. DATE ----------A2 19970624 JP 1995-347547 PΤ JP 09167584 19951218 In electron emitters, 1 side of elec. conductive films regarding AB electron emission parts is covered with metal films (e.g., refractory type), while the other with metal oxide films. The electron emitters have excellent emission characteristic and long life. Electron sources and display devices using the electron emitters are also described. Electrooptical imaging devices IT (elec. conductive films covered with metal oxides for electron emitters, electron sources and) IT Electric conductors (elec. conductive films covered with metal oxides for electron emitters, electron sources, and display devices) Refractory metal oxides IT Refractory metals RL: DEV (Device component use); USES (Uses) (elec. conductive films covered with metal oxides for electron emitters, electron sources, and display devices) IT Cathodes Electron emission (electron emitters, electron sources, and display devices) IT1304-28-5, Barium oxide (BaO), uses 1305-78-8, Calcium oxide (CaO), uses 1314-36-9, Yttrium oxide (Y2O3), uses 1312-81-8, Lanthanum oxide (La2O3) 12055-23-1, Hafnium oxide (HfO2) 13463-67-7, Titania, uses RL: DEV (Device component use); USES (Uses) (elec. conductive films covered with metal oxides for electron emitters, electron sources, and display devices) L29 ANSWER 9 OF 25 HCAPLUS COPYRIGHT 2002 ACS AN1997:211126 HCAPLUS DN 126:206544 TIElectron-emitting device, an electron source and image-forming apparatus using it, and their manufacture INShibata, Masaaki; Yamanobe, Masato; Tsukamoto, Takeo; Yamamoto, Keisuke; Arai, Yutaka PΑ Canon K. K., Japan

CODEN: EPXXDW Patent English LA FAN.CNT 1 KIND DATE APPLICATION NO. DATE PATENT NO. _____ -----EP 757371 A2 19970205 EP 1996-305645 19960731 PΙ EP 757371 A3 19970409 20001025 EP 757371 B1 R: DE, FR, GB, IT, NL JP 1996-197272 JP 09102267 A2 19970415 19960726 JP 3174999 B2 20010611 19970430 CN 1996-112123 19960801 CN 1148728 Α B1 20010206 US 6184610 US 1996-690964 19960801 CA 1996-2182647 19960802 AA 19970204 CA 2182647 AU 9660884 A1 19970206 AU 1996-60884 19960802 AU 711404 B2 19991014 PRAI JP 1995-216527 A 19950803 JP 1995-216542 A 19950803 JP 1995-216543 A 19950803 JP 1996-197272 A 19960726 JP 1996-197272 Α 19960726 AΒ An electron-emitting device comprises a pair of oppositely disposed device electrodes and an elec. conductive film elec. connecting the device electrodes and having an electron-emitting region formed as part of it. The elec. conductive film is partly or entirely covered by a metal oxide coating contg., as a principal ingredient, a metal oxide different from the material of the elec. conductive film, and having a higher m.p. and a lower work function than those of the principal ingredient of the elec. conductive film. The elec. conductive film also has a deposited layer comprising C, a C compd., or their mixt. IT Oxides (inorganic), processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (manuf. of electron emitters contg.) Electron sources TT Optical imaging devices (manuf. of electron emitters for) ΤT Cathodes (manuf. of electron emitters for electron sources and image-forming app.) TТ Television (manuf. of electron emitters for imaging devices for) 1304-28-5, Barium oxide (BaO), processes 1304-56-9, Beryllium oxide IT 1305-78-8, Calcium oxide (CaO), processes 1309-48-4, Magnesium oxide (MgO), processes 1312-81-8, Lanthanum oxide (La2O3) 1314-08-5, 1314-23-4, Zirconium oxide (ZrO2), processes Palladium oxide (PdO) 1314-35-8, Tungsten oxide (WO3), processes 1314-36-9, Yttrium oxide (Y2O3), processes 1332-37-2, Iron oxide, processes 1344-28-1, Aluminum oxide (Al2O3), processes 7440-02-0, Nickel, processes 7440-05-3, Palladium, processes 7440-06-4, Platinum, processes 7440-32-6, Titanium, processes 7440-47-3, Chromium, processes 7440-57-5, Gold, processes 7631-86-9, Silica, processes 12055-23-1, Hafnium oxide 13463-67-7, Titanium oxide (TiO2), processes 37300-04-2, Thorium oxide RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (manuf. of electron emitters contg.)

Serial No.:09/846,127

04/08/2002

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1996:393763 HCAPLUS
DN
    Detection method of silver halide photographic material
ΤI
    Shibata, Minoru; Kubota, Toshiharu; Egashira, Tetsutaro; Oda, Toshihiro;
IN
     Ito, Yoshimitsu
    Fuji Photo Film Co Ltd, Japan
PA
     Jpn. Kokai Tokkyo Koho, 10 pp.
SO
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                          APPLICATION NO. DATE
     ----- ----
                           -----
                                                          19940922
                     A2 19960412
                                          JP 1994-254660
PΙ
    JP 08095198
    A silver halide photog. material with Ag content .ltoreq.4.0 g/m2 composed
AB
    of a transparent support, a photosensitive Ag halide emulsion layer formed
     on the support, and a layer, which reflects IR and transmits visible
     radiation, contg. .gtoreq.1 metal oxide selected from
     In2O3, SnO2, ZnO, Al2O3, and TiO2, is detected with an IR sensor composed
     of an IR emitter and a receptor by detecting the redn. of IR
     transmittance while the photog. material is going through between the IR
     emitter and the receptor. The layer may be a primer formed
    between the support and the emulsion layer.
IT
    Electrophotography
     Infrared radiation
        (detection method of silver halide photog. material with IR sensor)
IT
    Optical detectors
        (IR, detection method of silver halide photog. material with IR sensor)
     7785-23-1, Silver bromide
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (detection method of silver halide photog. material with IR sensor)
     1312-43-2, Indium oxide (In2O3) 1314-13-2, Zinc oxide, uses
IT
     Alumina, uses 13463-67-7, Titania, uses
                                              18282-10-5, Tin oxide
             50926-11-9, ITO
     RL: TEM (Technical or engineered material use); USES (Uses)
        (in IR-reflective and visible radiation-transmit layer; detection
       method of silver halide photog. material with IR sensor)
    ANSWER 11 OF 25 HCAPLUS COPYRIGHT 2002 ACS
     1995:531138 HCAPLUS
AN
DN
     122:325874
ΤI
     Strongly-directed emission from microcavity structure in
     electroluminescent diodes with europium complex as an emitter
     Takada, Noriyuki; Tsutusi, Tetsuo; Saito, Shogo
ΑU
     Dep. Maters. Sci. and Technol., Kyushu Univ., Fukuoka, 816, Japan
CS
     Synth. Met. (1995), 71(1-3), 2099-100
SO
     CODEN: SYMEDZ; ISSN: 0379-6779
DT
     Journal
LΑ
    English
    Multilayer electroluminescent (EL) diodes with sharp red emission were
AB
     fabricated using an Eu complex as an emitter. Microcavity
     structures were introduced into the EL diodes with sharp red emission.
     The device structures were composed of pairs of TiO2/SiO2 dielec.
     reflector, In-Sn-oxide electrode, hole transport layer (HTL), Eu complex
     as an emission layer (EML), electron transport layer (ETL) and MgAg
     electrode. The dielec. reflector and the MgAG metal
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electrode constituted a planar microcavity. Sharply directed emission from Eu complex was obsd. when operated under d.c. drive voltage at room temp. In fabricating such EL diodes with optical microcavity made of

O

dielec. reflector and a metal mirror, the best total thickness of the org. layers depends on the stack structure of dielec. reflector. IT Electroluminescent devices (strongly-directed emission from microcavity structure in electroluminescent diodes with europium complex as emitter) 13463-67-7, Titanium oxide (TiO2), uses 50926-11-9, ITO IT 60676-86-0, Vitreous silica 65181-78-4 138372-67-5 RL: DEV (Device component use); USES (Uses) (strongly-directed emission from microcavity structure in electroluminescent diodes with europium complex as emitter) ANSWER 12 OF 25 HCAPLUS COPYRIGHT 2002 ACS L29 AN 1995:516262 HCAPLUS 122:251745 DN Manufacture of far-IR radiator sheet material TΙ IN Oikawa, Jusuke Shinnippon Seitetsu Kk, Japan PΑ SO Jpn. Kokai Tokkyo Koho, 4 pp. CODEN: JKXXAF Patent DΤ Japanese LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE PΙ JP 06299341 A2 19941025 JP 1993-104906 19930408 AB A far-IR radiating sheet material, suited for use in heaters, is made by forming a Ti oxide thin film on a metal plate by ion plating in an Ar atm. with an O2 partial pressure 0.4-0.6 mtorr. Infrared sources ΙT (far-, manuf. of far-IR emitter sheet) 12137-20-1, Titanium monoxide 13463-67-7, Titania, uses IT RL: TEM (Technical or engineered material use); USES (Uses) (manuf. of far-IR emitter sheet) ANSWER 13 OF 25 HCAPLUS COPYRIGHT 2002 ACS 1994:566729 HCAPLUS AN121:166729 DNMolecular-Level Electron Transfer and Excited State Assemblies on Surfaces ΤI of Metal Oxides and Glass Meyer, Thomas J.; Meyer, Gerald J.; Pfennig, Brian W.; Schoonover, Jon R.; ΑU Timpson, Cliff J.; Wall, Jennifer F.; Kobusch, Claus; Chen, Xiaohong; Peek, Brian M.; et al. Department of Chemistry, University of North Carolina, Chapel Hill, NC, CS 27599-3290, USA Inorg. Chem. (1994), 33(18), 3952-64 SO CODEN: INOCAJ; ISSN: 0020-1669 DΤ Journal English A general procedure is described for the attachment to antimony-doped tin dioxide (SnO2:Sb), tin-doped indium oxide(In2O3:Sn), or glass surfaces of mols. with known electron transfer or excited state properties, e.g.

LA AB [Ru(bpy) 2 (4,4-(CO2H) 2bpy)] (PF6) 2 (bpy = 2,2'-bipyridine; 4,4-(CO2H) 2bpy = 2,2'-bipyridine; 4,4'-(CO2H) 2bpy = 2,2'-bipyridine; 4,4'-(CO2H) 2bpy = 2,2'-bipyridine; 4,4'-(CO2H) 2bpy = 2,2'-bipyridine; 4,4'-(CO2H) 2bpy = 2,2'-bipyridine; 4,2'-bipyridine; 4,4'-(CO2H) 2bpy = 2,2'-bipyridine; 4,4'-(CO2H) 2bpyridine; 4,4'-(CO2H4,4-dicarboxy-2,2'-bipyridine), based on the interaction between surface hydroxyls and carboxylic acid groups. Integrations of cyclic voltammetric waveforms on the metal oxide electrodes give max. surface coverages of .GAMMA. .apprx. 1 .times.10-10 mol/cm2 for the ruthenium complex, which corresponds to a monolayer coverage. At. force

microscope (AFM) measurements reveal that the metal oxide surfaces are highly roughened with root mean square roughnesses in the range 4-6.5 nm for tin oxide. The smaller orgs., N-methyl-N-viologenpropanoic acid bis(hexafluorophosphate), [MV-CO2H] (PF6)2, and 10H-phenothiazine-10-propanoic acid, PTZ-CO2H, display similar surface coverages. Resonance Raman measurements on surfaces contq. the ruthenium complex imply that attachment to SnO2, In203, and TiO2 is via an ester bond. For SiO2, two modes of binding are suggested, a majority involving a chelating carboxylato link and a minority, ester formation. Binding consts. for surface attachment were measured in CH2Cl2 at 298 K by equilibration, which gave K = 8 .times. 104 M-1 on both SnO2:Sb and In2O3:Sn. Surface mol. assemblies have been prepd. contg. [Ru(bpy)2(4,4-(CO2H)2bpy)](PF6)2 and [Os(bpy)2(4,4-(CO2H)2 bpy)](PF6)2, [MV-CO2H](PF6)2, and PTZ-CO2H. In these assemblies, sep. waves are obsd. for the different redox couples at potentials near those found for surfaces contg. only a single component. Emission decay of the metal-to-ligand charge transfer (MLCT) excited state of [Ru(bpy)2(4,4'-(CO2H)2bpy)] (PF6)2 attached to the glass backings of metal oxide electrodes or to glass slides was found to be nonexponential with av. lifetimes that varied from < 5 to 550 ns with CH2Cl2 in the external soln. The data could be satisfactorily fit to the Williams-Watts (Kohlrausch) distribution function. Studies at varying surface coverages revealed that the av. lifetime of emission decay (.ltbbrac..tau..rtbbrac.) depends on the extent of surface coverage and increases as surface coverage decreases. There is evidence for excited state-ground state interactions by a red-shift in the emission max. as the extent of surface coverage is increased. Emission decay remains nonexponential even on surfaces that are lightly covered. The emission was almost completely quenched on the semiconductor surfaces, with .ltbbrac..tau..rtbbrac. < 5 ns. The bound Ru(II) emitters on glass were quenched by electron or energy transfer to the coattached quenchers [MV-CO2H] (PF6)2, PTZ-CO2H, or [Os(bpy)2(4,4'-(CO2H)2bpy)] (PF6)2, suggesting that lateral electron and energy transfer can occur across the surface. Surface lifetime quenching also occurred in the presence of added 10-methyl-10-phenothiazine in the external CH2Cl2 soln. The kinetics of lifetime quenching did not follow Stern-Volmer kinetics but could be fit to a model in which there are both quenchable and unquenchable sites on the same surface.

ΤТ Electrodes

(metal oxide, mol.-level electron transfer and excited state assemblies on surfaces of)

IT Energy transfer

Kinetics of photolysis

Redox reaction

(of ruthenium complexes on metal oxide surfaces,

electron transfer in)

TT Luminescence

Ultraviolet and visible spectra

(of ruthenium complexes on metal oxide surfaces,

for mol.-level assemblies with well defined redox properties)

IT Photochemistry

> (photophysics and, of ruthenium complexes on metal oxide surfaces, electron transfer in)

IT Adsorbed substances

> (monolayer, of ruthenium complexes on metal oxide surfaces, electron transfer in)

Electron exchange and Charge transfer TT

(photochem., of ruthenium complex mol. assemblies on metal oxide surfaces)

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Raman spectra
IT
        (resonance, of ruthenium complexes on metal oxide
        surfaces, electron transfer in)
    18282-10-5, Tin dioxide
IT
    RL: USES (Uses)
        (antimony-doped, mol.-level electron transfer and excited state
       assemblies on surfaces of)
    7440-31-5, Tin, uses
IΤ
    RL: USES (Uses)
        (indium oxide electrode doped with, mol.-level electron transfer and
        excited state assemblies on surfaces of)
     1207-72-3, 10-Methylphenothiazine
IT
    RL: USES (Uses)
        (luminescence quenching of ruthenium complexes by, on metal
        oxide surfaces)
     7631-86-9, Silica, uses 13463-67-7, Titanium dioxide, uses
IT
    RL: USES (Uses)
        (mol.-level electron transfer and excited state assemblies on surfaces
       of)
     362-03-8, 10H-Phenothiazine-10-propanoic acid 62207-96-9 68264-89-1
TΤ
     92984-72-0 145205-16-9 157473-46-6 157473-47-7
    RL: USES (Uses)
        (surface attachment and photophys. properties of, as mol.-level
        assemblies with well defined redox properties)
ΙT
    7440-36-0, Antimony, uses
    RL: USES (Uses)
        (tin dioxide electrode doped with, mol.-level electron transfer and
        excited state assemblies on surfaces of)
IT
    1312-43-2, Indium oxide (In2O3)
    RL: USES (Uses)
        (tin-doped, mol.-level electron transfer and excited state assemblies
       on surfaces of)
    ANSWER 14 OF 25 HCAPLUS COPYRIGHT 2002 ACS
L29
AN
    1993:615789 HCAPLUS
    119:215789
DN
TI
    Insulating channel dielectric-base transistor
IN
    Tamura, Yasutaka
PΑ
    Fujitsu Ltd, Japan
SO
    Jpn. Kokai Tokkyo Koho, 8 pp.
    CODEN: JKXXAF
DT
    Patent
LA
    Japanese
FAN.CNT 1
                     KIND DATE
    PATENT NO.
                                          APPLICATION NO. DATE
     _____
                                           -----
PΤ
    JP 05198854
                     A2
                           19930806
                                          JP 1992-9026
                                                           19920122
    The transistor comprises a base region having a high dielec. const., an
AR
    emitter electrode on 1 side of the base region through a 1st
    barrier layer having a dielec. const. lower than that of the base region,
    an collector electrode through a 2nd barrier layer of the same type, and a
    base electrode on the other side through a 3rd barrier layer with height
    and thickness to prevent carrier tunneling. The 3rd barrier layer may
    have the dielec. const. higher than those of the 1st and 2nd barrier
    layers. The transistor may comprise an insulating substrate successively
    covered with the dielec. base region layer, the 3rd barrier layer and the
    base electrode. The base electrode may be made of an impurity-doped or
    quasi-stoichiometric base region material. The emitter,
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collector, and/or base electrode may consist of metal or

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oxide superconductor. The base region may consist of oxide contg.
    Sr, Ti, K, Ta, Sn, Zr, or Nb. The base region may consist of KTal-xNbxO3,
    having controlled temp. of the max. dielec. const. with the Nb content.
IT
    Transistors
        (dielec.-base, with tunneling barrier layer)
    Electric insulators and Dielectrics
IT
        (transistor base from)
    7631-86-9, Silica, uses
                              12003-86-0, Aluminum yttrium oxide (AlYO3)
IT
    13463-67-7, Titanium dioxide, uses
    RL: USES (Uses)
        (dielec.-base transistor carrier tunneling barrier layer)
ΙT
    7440-25-7, Tantalum, uses 107539-20-8, Barium copper yttrium oxide
    RL: USES (Uses)
        (dielec.-base transistor electrode)
     7440-03-1, Niobium, uses
IT
    RL: USES (Uses)
        (strontium titanate doped with, for dielec.-base transistor electrode)
     12060-59-2, Strontium titanium oxide (SrTiO3) 12710-39-3, Niobium
IT
    potassium tantalum oxide
    RL: USES (Uses)
        (transistor dielec.-base from, with carrier tunneling barrier layer)
L29
    ANSWER 15 OF 25 HCAPLUS COPYRIGHT 2002 ACS
     1991:473866 HCAPLUS
AN
DN
    115:73866
TΙ
    Antibacterial, deodorant, far-IR-radiating antistatic paper
IN
    Yoshizawa, Noriyasu
PΑ
    Ain Corp., Ltd., Japan
    Jpn. Kokai Tokkyo Koho, 9 pp.
SO
    CODEN: JKXXAF
    Patent
DT
LA
    Japanese
FAN.CNT 1
                    KIND DATE
    PATENT NO.
                                          APPLICATION NO. DATE
     -----
                                          -----
    JP 03069695
                     A2
                           19910326
                                          JP 1989-222914 19890831
PΤ
PRAI JP 1989-114104
                          19890509
    The title paper and paperboard are manufd. by coating paper base with a
     ceramic compn. contg. far-IR-radiating and antistatic metal
     oxide binders, synthetic polymers, and antibacterial and
     gas-scavenging inorg. fillers. Typical metal oxides
     are SiO2, Al oxide, and TiO2; synthetic polymers are modified
    polypropylenes; and antibacterial and gas-scavenging fillers contain metal
     ions e.g. Cu and Ag ions and zeolites, etc.
IT
     Paper
        (antibacterial, antistatic, deodoring and far-IR-emitting ceramics
        coatings on)
IT
    Oxides, uses and miscellaneous
     Zeolites, uses and miscellaneous
     RL: USES (Uses)
        (ceramic coatings contg., on paper, for antibacterial, antistatic,
       deodoring and far IR-emitting properties)
    Bactericides, Disinfectants, and Antiseptics
ΙT
        (in ceramic coatings on paper, heavy metal ions as)
IT
     Antistatic agents
        (in ceramic coatings on paper, metal oxides as)
IT
    Deodorants
        (in ceramic coatings on paper, zeolites as)
IT
     Binding materials
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(inorg. compds., in ceramic coatings on paper, metal
        oxides as)
     Coating materials
IT
        (ceramic, on paper, contg. far IR emitters, binders,
        antistatic and antibacterial agents and deodorants)
     14701-21-4, Silver ion, uses and miscellaneous 15158-11-9, uses and
IT
     miscellaneous
    RL: USES (Uses)
        (ceramic coatings contg., on paper bearing antibacterial, antistatic,
        deodoring and far IR-emitting properties)
     1344-28-1, Aluminum oxide, uses and miscellaneous
                                                         7631-86-9, Silicon
IT
    oxide, uses and miscellaneous 9003-07-0D, Polypropylene, derivs.
     13463-67-7, Titanium oxide, uses and miscellaneous
    RL: USES (Uses)
        (ceramic coatings contg., on paper, for antibacterial, antistatic,
        deodoring and far IR-emitting properties)
L29 ANSWER 16 OF 25 HCAPLUS COPYRIGHT 2002 ACS
    1991:473844 HCAPLUS
AN
    115:73844
DN
ΤI
    Antibacterial, antistatic, deodoring and far-IR-emitting coatings on wood
IN
    Yoshizawa, Noriyasu
PΑ
    Ain Corp., Ltd., Japan
    Jpn. Kokai Tokkyo Koho, 9 pp.
SO
     CODEN: JKXXAF
DT
    Patent
LA
    Japanese
FAN.CNT 1
                     KIND DATE
     PATENT NO.
                                           APPLICATION NO. DATE
                     ----
    JP 03073304
                      A2
                            19910328
                                           JP 1989-222918
                                                            19890831
PRAI JP 1989-114108
                            19890509
    The title coatings comprise far-IR-emitting and antistatic metal
     oxide binders, e.g. SiO2, Al2O3, and TiO2; synthetic polymer
    binders, e.g. modified polypropylene; and antibacterial and gas-scavenging
     inorg. fillers, e.g. from Cu and Ag ions and zeolites.
    Wood
IT
        (antibacterial and antistatic and deodorizing and far-IR-emitting
        ceramic coatings on)
TΤ
    Oxides, uses and miscellaneous
     Zeolites, uses and miscellaneous
     RL: USES (Uses)
        (ceramic coatings contg., on wood, for antibacterial, antistatic,
        deodorizing and far IR-emitting properties)
    Bactericides, Disinfectants, and Antiseptics
IT
        (in ceramic coatings on wood, heavy metal ions as)
ΙT
    Antistatic agents
        (in ceramic coatings on wood, metal oxides as)
IT
    Deodorants
        (in ceramic coatings on wood, zeolites as)
TT
    Binding materials
        (inorg. compds., in ceramic coatings on wood, metal
       oxides as)
ΙT
    Coating materials
        (ceramic, on wood, contg. far IR emitters, binders,
        antistatic and antibacterial compns. and deodorants)
     Ceramic materials and wares
IT
        (coatings, on wood, contg. far IR emitters, binders,
       antistatic and antibacterial compns. and deodorants)
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1344-28-1, Aluminum oxide, uses and miscellaneous
                                                        7631-86-9, Silicon
    oxide, uses and miscellaneous 9003-07-0D, Polypropylene, derivs.
    13463-67-7, Titanium oxide, uses and miscellaneous 14701-21-4,
    Silver ion, uses and miscellaneous 15158-11-9, uses and miscellaneous
    RL: USES (Uses)
        (ceramic coatings contg., on wood, for antibacterial, antistatic,
        deodorizing and far IR-emitting properties)
    ANSWER 17 OF 25 HCAPLUS COPYRIGHT 2002 ACS
L29
    1991:446341 HCAPLUS
AN
    115:46341
DN
ΤI
    Manufacture of tobacco filters with deodorizing, antibiotic, far-infrared
    radiation and antistatic properties
    Yoshizawa, Noryasu
IN
    Ain Corp., Ltd., Japan
PΑ
     Jpn. Kokai Tokkyo Koho, 9 pp.
SO
     CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
    PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
     19910409
PΙ
     JP 03083571 A2
                                           JP 1989-220477
                                                             19890829
     The filter can be made by coating a filter which consists of various
AB
     fibers, with a ceramic coating agent which consists of: (1) adhesive,
     far-IR irradg. and antistatic metal oxide (e.g. SiO2,
     Al203, TiO2) (2) adhesive resin (e.g. modified polypropylene), and (3)
     antibiotic, gas-absorbing, and gas-decompg. inorg. filler (e.g. metal ion,
    zeolite). The coating can be done by spraying, dipping, rolling, etc. Thus, a coating agent powder consisting of SiO2 (as adhesive and far-IR
     irradg. agent) , Al203 (as adhesive and antistatic agent), TiO2 (as
     adhesive and deodorizing agent, modified polypropylene (As adhesive), Cu
     ion or Ag+ (as antibiotic and gas-decompg. agent), zeolite
     (Ca.cntdot.2Al203.cntdot.5SiO2; as gas-absorbent) was dissolved in alc.
     and sprayed to acetate fiber filter (diagram of app. for coating is
     given), the antibiotic and deodorizing effects were demonstrated.
IT
     Zeolites, biological studies
    RL: BIOL (Biological study)
        (as smoke absorbent in manuf. of cigarette filter)
     Tobacco smoke and smoking
IT
        (filters for, antimicrobial and far-IR irradg. materials in)
TT
    Antistatic agents
    Deodorants
        (tobacco filter manuf. with)
TT
    Coating materials
        (ceramic, tobacco filter manuf. with)
TT
    Ceramic materials and wares
        (coatings, tobacco filter manuf. with)
     Infrared radiation
TT
        (far-, emitters of, in manuf. of cigarette filter)
IT
     7440-22-4, Silver, biological studies
    RL: BIOL (Biological study)
        (as antibiotic and gas-decompg. agent in manuf. of cigarette filter)
     1344-28-1, Aluminum oxide, biological studies 7631-86-9, Silicon
TT
     dioxide, biological studies 13463-67-7, Titanium oxide,
     biological studies
    RL: BIOL (Biological study)
        (tobacco filter contg.)
     7440-50-8, Copper, biological studies 9003-07-0, Polypropylene
IT
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PATENT NO.

JP 02022480

ΡI

KIND DATE

19900125

A2

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RL: BIOL (Biological study)
        (tobacco filter manuf. with)
    1335-30-4
IT
     RL: BIOL (Biological study)
        (zeolites, as smoke absorbent in manuf. of cigarette filter)
    ANSWER 18 OF 25 HCAPLUS COPYRIGHT 2002 ACS
T.29
    1991:220251 HCAPLUS
AN
DN
     114:220251
     Use of phosphate fluxes for the preparation of homogeneous x-ray
ΤI
     Karmanov, V. I.; Slobodyanik, N. S.; Nagornyi, P. G.; Ryabushko, O. P.
ΑU
     E. O. Paton Inst. Electr. Welding, Kiev, USSR
CS
     Zh. Anal. Khim. (1991), 46(1), 121-6
     CODEN: ZAKHA8; ISSN: 0044-4502
DT
     Journal
     Russian
LA
     Potassium and sodium tetraborates have been commonly used as fluxes in
ΑB
     sample prepn. for x-ray fluorescence anal. Because of the low soly. this
     method can not be successfully applied to the detn. of polyvalent
     metal oxides. Phosphate flux is suggested and studied
     in this paper for prepg. homogeneous glassy x-ray emitters of
     polyvalent metal oxides. As an example, the method
     was used for the prepn. of synthetic ref. samples for x-ray fluorescence
     detn. of Al203, Fe203, Si02, Ti02, and ZrO2 in rutile samples. The mixts.
     of NaPO3 and Na4P2O7 with different ratios were tested for obtaining best
     soly. of rutile. Three industrial rutile samples were analyzed. The
     results compared well with those of chem. anal.
     Fluxes
IT
        (phosphates, for polyvalent metal detn. by x-ray fluorescence)
TΤ
     X-ray analysis
        (fluorescence, polyvalent metal detn. by, phosphate flux in)
     7429-90-5, Aluminum oxide, analysis 7439-89-6, Iron oxide (Fe2O3), analysis 7440-21-3, Silicon dioxide, analysis 7440-32-6, Titanium
TT
                         7440-67-7, Zirconium dioxide, analysis
     dioxide, analysis
     RL: ANT (Analyte); ANST (Analytical study)
        (detn. of, in rutile by x-ray fluorescence using phosphate flux)
     7722-88-5, Tetrasodium pyrophosphate 10361-03-2, Monosodium phosphate
IT
     RL: ANST (Analytical study)
        (in flux for polyvalent metal oxide detn. by x-ray
        fluorescence)
     1317-80-2, Rutile
IT
     RL: ANST (Analytical study)
        (polyvalent metal detn. in, by x-ray fluorescence using phosphate flux)
     ANSWER 19 OF 25 HCAPLUS COPYRIGHT 2002 ACS
L29
     1991:8335 HCAPLUS
AN
DN
     114:8335
     Coated metal plate with good workability and far-IR emission properties
ΤI
     Yamazaki, Takao; Kanai, Hiroshi; Oka, Joji
IN
PΑ
     Nippon Steel Corp., Japan
     Jpn. Kokai Tokkyo Koho, 6 pp.
SO
     CODEN: JKXXAF
DT
     Patent
LA
     Japanese
FAN.CNT 1
```

APPLICATION NO. DATE

19880712

JP 1988-171817

```
The title plate is formed by adhering an org. polymer film contg. 5-95%
    far-IR emitter to a metal plate. A typical far-IR
    emitter was obtained by sintering a mixt. of MnO2 50, Fe2O3 20,
    CuO 10, CoO 10, and ZrO2 10% at 1200.degree.. A polyethylene film contg.
     60% far-IR emitter was adhered on a tinplate using a polymer
    adhesive by hot pressing at 180.degree..
    Coating materials
        (far-IR emitter-contg., for tinplates, with good workability)
ΙT
    Infrared sources
        (far-, metal oxides, coatings contg., on timplate)
     9002-88-4, Polyethylene
ΙT
    RL: TEM (Technical or engineered material use); USES (Uses)
        (coatings, contg. far-IR emitters, for timplate)
     1304-28-5, Barium oxide, uses and miscellaneous
                                                    1305-78-8, Calcium
TТ
    oxide, uses and miscellaneous 1307-96-6, Cobalt oxide (CoO), uses and
                   1308-38-9, Chromium oxide (Cr2O3), uses and miscellaneous
    miscellaneous
     1309-37-1, Ferric oxide, uses and miscellaneous
                                                    1309-48-4, Magnesium
    oxide (MqO), uses and miscellaneous 1313-13-9, Manganese dioxide, uses
                        1313-99-1, Nickel oxide, uses and miscellaneous
     and miscellaneous
     1314-23-4, Zirconium oxide, uses and miscellaneous 1317-38-0, Cupric
    oxide, uses and miscellaneous 1344-28-1, Aluminum oxide, uses and
    miscellaneous
                   7631-86-9, Silica, uses and miscellaneous
                                                               12057-24-8,
    Lithium oxide, uses and miscellaneous 13463-67-7, Titania, uses
     and miscellaneous
    RL: USES (Uses)
        (far-IR emitters contg., in polymeric coatings, on tinplate
       with good workability)
    ANSWER 20 OF 25 HCAPLUS COPYRIGHT 2002 ACS
L29
     1988:612889 HCAPLUS
AN
     109:212889
DN
TI
     An efficient and durable aluminum or aluminum alloy far-IR radiator
     Ishida, Shinichi; Yamada, Kikuo
IN
PA
     Nippon Aluminium Mfg. Co., Ltd., Japan
     Jpn. Kokai Tokkyo Koho, 6 pp.
SO
     CODEN: JKXXAF
DT
     Patent
LA
    Japanese
FAN.CNT 1
                    KIND DATE
     PATENT NO.
                                          APPLICATION NO. DATE
     ----- ----
                                          ______
                         19880617
                                          JP 1986-292121 19861208
PΙ
     JP 63145797
                     A2
     The radiator comprises an anodic oxide film including far-IR emitting
AB
     materials. Optionally, the far-IR emitting material may comprise C, or an
     oxide of Fe, Cr, Ni, Co, Ti, Sn, Ag, Pb, Au, Mg, Mn, Zn, etc. The
     radiator is useful as a heating component.
    Heating systems and Heaters
IT
        (aluminum far-IR emitters)
     Transition metal oxides
IT
     RL: USES (Uses)
        (far-IR radiators from anodized aluminum with)
IT
     Infrared sources
        (far-, anodized aluminum)
IT
     7429-90-5, Aluminum, uses and miscellaneous 11121-92-9
```

1309-48-4, Magnesium oxide, uses and miscellaneous TT 1314-13-2, Zinc oxide, uses and miscellaneous 1332-29-2, Tin oxide 1332-37-2, Iron oxide (unspecified), uses and miscellaneous 1335-25-7, Lead oxide

RL: USES (Uses)

(far-IR radiators from anodized)

- ANSWER 22 OF 25 HCAPLUS COPYRIGHT 2002 ACS L29
- 1985:493358 HCAPLUS AN
- DN 103:93358
- X-ray photoelectron diffraction (XPED) studies on metal ΤI oxide surfaces. (I). Analysis of the XPED patterns from titanium dioxide (001) and .alpha.-aluminum oxide (0001) by the single scattering calculation
- Tamura, Koji; Owari, Masanori; Kudo, Masahiro; Nihei, Yoshimasa ΑU
- CS
- Inst. Ind. Sci., Univ. Tokyo, Tokyo, 106, Japan Bull. Chem. Soc. Jpn. (1985), 58(7), 1873-8 SO CODEN: BCSJA8; ISSN: 0009-2673
- Journal DT
- English LA
- X-ray photoelectron diffraction (XPED) measurements were applied to the AΒ (001) surface of TiO2 rutile and the (0001) surface of .alpha.-Al2O3 to examine the applicability of XPED to metal oxides. Obtained XPED patterns were compared with the results of theor. calcns. based on a single scattering model. The calcd. patterns from both oxides agreed with the exptl. patterns. The 4 O atoms in the unit cell of TiO2 make different contributions to the XPED pattern which reflects different at. environments. The same calcn. was also made on Ti in TiO2, and on Al and O in Al2O3. This clarified the contributions of nonequivalent sites of emitter atoms to the main peaks in the exptl. XPED patterns. In comparison with the crystal structures, some peaks could be attributed to some specific emitters or specific emitter -scatterer pairs.
- ΙT Surface structure

STIC-EIC 2800 CP4-9C18

```
(detn. of, of metal oxides, x-ray photoelectron
        diffraction in)
IT Photoelectric emission
        (x-ray, in surface structure detn.)
IT
     7782-44-7, properties
     RL: PRP (Properties)
        (coordination of, in alumina and titania crystals, x-ray photoelectron
        diffraction in detn. of)
IT
     1344-28-1, properties 13463-67-7, properties
     RL: PRP (Properties)
        (surface structure detn. of, x-ray photoelectron diffractometry in)
    ANSWER 23 OF 25 HCAPLUS COPYRIGHT 2002 ACS
L29
AN
     1978:92160 HCAPLUS
     88:92160
DN
    Thermionic converter performance with oxide collectors
ΤI
     Lieb, D.; Goodale, D.; Briere, T.; Balestra, C.
ΑU
     Thermo Electron Corp., Waltham, Mass., USA
CS
     Proc. - Intersoc. Energy Convers. Eng. Conf. (1977), 12, Vol. 2, 1555-62
SO
     CODEN: PIECDE
DT
     Journal
    English
LA
AΒ
     Thermionic converters using various metal-oxide
     collector surfaces were fabricated and tested. Work function and power
     output data are presented and evaluated. Oxides of Ba, Sr, Zn, W, and Ti
     were incorporated into a variable-spacing converter. W oxide gave the
     highest converter performance and simultaneously furnished O for the
     emitter. Oxygenated emitters operate at reduced Cs
     pressure with an increase in electrode spacing. Cs penetration of the
     W-oxide layer was detd. with possible formation of a Cs tungstate bronze.
     Ti oxide showed high performance but did not furnish O for the
     emitter. SrO in the form of a sprayed layer dissocd. in the
     presence of Cs. Sprayed coatings of BaO and ZnO produced collector work
     functions of .apprx.1.3 eV but had excessive series resistance. LaB6 in
     combination with O introduced through a Ag tube and Cs produced a
     low-work-function collector and better than av. performance.
IT
    Work function
        (of oxide collectors, thermionic-converter performance and)
IT
     Thermionic energy converters
        (with oxide collectors, performance of)
IT
     1304-28-5, properties
                            1314-11-0, properties
                                                     1314-13-2, properties
                            12008-21-8 13463-67-7, properties
     1314-35-8, properties
     RL: PRP (Properties)
        (thermionic converter with collector surfaces of, properties of)
    ANSWER 24 OF 25 HCAPLUS COPYRIGHT 2002 ACS
L29
     1973:10091 HCAPLUS
AN
     78:10091
ĎΝ
ΤI
     Applying a multilayer antireflection coating to a substrate
     Louderback, Anthony W.; Zook, Morris A., Jr.
IN
     U.S., 7 pp. Division of U.S. 3,604,784 (CA 75; 146130n).
SO
     CODEN: USXXAM
DT
     Patent
LA
    English
FAN.CNT 1
                    KIND DATE
     PATENT NO.
                                          APPLICATION NO. DATE
     -----, ---- , ----
PΙ
    US 3695910
                      Α
                            19721003
                                          US 1970-50010
                                                            19700528
AΒ
     A non-absorbing colorless, multilayer antireflective coating is deposited
```

TT

IT

TT

IT

ΙT

ANDN

TIIN

PA

SO

DT

LA

PΙ

AB

on a substrate having light reflecting properties by placing the substrate in a vacuum chamber contg. coating materials of metal oxides and metal fluorides a source of O and an electron beam emitter to effect evapn. of the materials. The chamber is evacuated to .ltoreq.3 .times. 10-5 torr, 1st layer of metal oxide is evapd. on the substrate then a mixt. of 2 metal oxides is evapd. on the 1st layer. A source of O is directed at the electron beam during evapn. of the 2nd layer to increase the pressure to a max. of 2.5 .times. 10-4 torr then a metal fluoride is evapd. on the 2nd layer. For example the substrates was placed in a holding device, coating materials were placed in receptacles and the system was evacuated to 3 .times. 10-5 torr. The substrate was heated to 500.degree.F. Fused Al2O3 (60 mesh) was evapd. to give a 1st layer. The middle layer was deposited from a mixt. of elec. fused Al2O3 and a powd. mixt. of TiO, Ti203, and Ti02. The final layer of MgF2 was deposited with an electron beam at 3 .times. 10-5 torr. Fluorides, uses and miscellaneous Oxides, uses and miscellaneous RL: USES (Uses) (in antireflection coating) Electron beam, chemical and physical effects (in multilayer antireflection coating of substrates) Coating process (multilayer antireflection substrate application) 7782-44-7, uses and miscellaneous RL: USES (Uses) (in antireflection coatiing application) 1344-28-1, uses and miscellaneous 1344-54-3 RL: USES (Uses) (in antireflection multilayer coating of substrates) 7783-40-6 12137-20-1 13463-67-7, uses and miscellaneous RL: PRP (Properties) (in antireflection multilayer coating of substrates) ANSWER 25 OF 25 HCAPLUS COPYRIGHT 2002 ACS 1967:495140 HCAPLUS 67:95140 Narrow-band emitter devices Schmidlin, Frederick W. GTC Corp. U.S., 10 pp. CODEN: USXXAM Patent English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ----------US 3310685 19670321 US 19630503 These tunnel emission cathodes and tunnel-emission amplifiers have narrow band emitters with low noise. In the tunnel cathode a high emission efficiency is achieved with an emitter material such as an n-type semiconductor of TiO, Ti203, or VO3 with a conduction band contg. electrons whose spread and energies extend over a narrow range. The emitter material is bonded to 1 side of a thin dielec. material such as polymd. siloxane which serves as the tunnel barrier. the opposite side of the siloxane is bonded a thin metal layer which serves as the base. The exposed surface of the base may be treated to

lower the vacuum work function. In the tunnel emission amplifier, the

ΙT

IT

TΤ

IT

IT

IT

heavily doped n-type semiconductor, the dielec., and the metal are bonded as in the tunnel cathode. Then a layer of an intrinsic semiconductor or a dielec. materials is bonded on the opposite side of the base to serve as the sluice. On the opposite side of the sluice a metal layer is bonded to serve as a collector. The sluice can be doped to warp its conduction band to obtain an improved transmission efficiency. These devices are operated with emitter-base voltages just large enough to raise the narrow band of electrons in the emitter to an energy level slightly above that required for an electron to surmount the base-vacuum or base-sluice work function. The amt. of scattering of the tunneled electrons is min. and the emission efficiency is a max. Siloxanes, uses and miscellaneous RL: USES (Uses) (electron-tunnel emission amplifiers and cathodes contg.) Electron emission (tunnel, amplifiers and cathodes based on narrow-band) Cathodes (tunnel-emission, oxide) Electric amplifiers (tunnel-emission, solid-state narrow-band) Potential barriers (tunneling through, in emitter devices) 1314-34-7 **1344-54-3 12137-20-1** RL: USES (Uses) (electron-tunnel emission amplifiers and cathodes contq.)

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L49 ANSWER 1 OF 11 HCAPLUS COPYRIGHT 2002 ACS
    2001:452643 HCAPLUS
AN
DN
    Electron-emitting material with low evaporation during discharge and
     resistant to ion sputtering and its electrode and electric discharge tube
    Takahashi, Makoto; Hamada, Munemitsu; Matsuoka, Hiroshi; Takeishi, Akira;
    Yodokawa, Masatada
PΑ
    TDK Corporation, Japan
     Jpn. Kokai Tokkyo Koho, 9 pp.
SO
     CODEN: JKXXAF
DT
    Patent
    Japanese
LΑ
FAN.CNT 1
                                           APPLICATION NO. DATE
                     KIND DATE
    PATENT NO.
     _____
     JP 2001167687
                     A2 20010622
                                          JP 1999-346967 19991206
PΙ
AB
     An electron-emitting material contains a 1st metal component selected from
     Ba, Sr and/or Ca and a 2nd metal component selected from Ta, Zr, Nb, Ti
     and/or Hf; when the 1st and the 2nd components are shown as MI and MII,
     resp., the material contains a mixed oxide involving MI5MII4015-type
     crystal as the base component. The material may contain a metal M (M .gtoreq.1 selected from Mg, Sc, Y, lanthanoid, V, Cr, Mo, W, Fe, Ni, Al).
     Preferably, the material contains .gtoreq.1 of a crystal type shown as
    MI4MII209, MIMII206, MI6MIIMII4018, MI7MII6022, and MIMII03. The elec.
     discharge tube assembled with an electrode of the material has suppressed
    blackening on the tube wall and hence has long service life.
IT
     Cathodes
     Electric discharge devices
        (electron-emitting material with low evapn. during discharge and
        resistant to ion sputtering and its electrode and elec. discharge tube)
TT
     345232-64-6P, Barium tantalum zirconium oxide (Ba5Ta3.92Zr0.08015)
     345232-66-8P, Barium tantalum zirconium oxide (Ba5Ta3.8Zr0.2015)
     345232-67-9P, Barium tantalum zirconium oxide (Ba5Ta3.6Zr0.4015)
     345232-68-0P, Barium tantalum zirconium oxide (Ba5Ta2.8Zr1.2015)
     345232-69-1P, Barium tantalum zirconium oxide (Ba5Ta2Zr2O15)
     345232-70-4P, Barium tantalum zirconium oxide (Ba5Tal.2Zr2.8015)
     345232-71-5P, Barium tantalum zirconium oxide (Ba5Ta0.4Zr3.6O15)
     345232-72-6P, Barium hafnium tantalum oxide (Ba5Hf2Ta2O15) 345232-73-7P,
     Barium niobium zirconium oxide (Ba5Nb2Zr2O15) 345232-74-8P, Barium
     hafnium zirconium oxide (Ba5Hf2Zr2O15) 345232-75-9P, Barium hafnium
     niobium oxide (Ba5Hf2Nb2O15)
                                  345232-76-0P, Barium hafnium
                                    345232-77-1P, Barium
     titanium oxide (Ba5Hf2Ti2O15)
     niobium tantalum zirconium oxide (Ba5Nb1.2Ta1.6Zr1.2O15)
                                                                345232-78-2P,
     Barium tantalum titanium zirconium oxide
     (Ba5Tal.6Til.2Zrl.2O15) 345232-79-3P, Barium hafnium tantalum zirconium
     oxide (Ba5Hf1.2Ta1.6Zr1.2O15)
                                    345232-80-6P, Barium hafnium niobium
     tantalum oxide (Ba5Hf1.2Nb1.2Ta1.6015)
                                              345232-81-7P, Barium hafnium
     tantalum titanium oxide (Ba5Hf1.2Ta1.6Ti1.2015)
     345232-82-8P, Barium niobium titanium zirconium oxide
     (Ba5Nbl.2Til.2Zrl.6O15) 345232-83-9P, Barium hafnium niobium zirconium
     oxide (Ba5Hf1.2Nb1.2Zr1.6O15)
                                    345232-84-0P, Barium hafnium
     titanium zirconium oxide (Ba5Hf1.2Ti1.2Zr1.6O15)
     345232-85-1P, Barium hafnium niobium titanium oxide
     (Ba5Hf1.2Nb1.6Ti1.2015)
                             345232-86-2P
L49 ANSWER 2 OF 11 HCAPLUS COPYRIGHT 2002 ACS
AN
     1999:165906 HCAPLUS
     130:290017
DN
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- Photoelectric work functions of metal oxide films and emission characteristics of molybdenum emitter tips with oxide coatings
- Bernhard, J. M.; Rouse, A.; Sosa, E. D.; Golden, D. E.; Chalamala, B. R.; ΑU Aggarwal, S.; Gnade, B. E.; Ramesh, R.
- Department of Physics, University of North Texas, Denton, TX, 76203, USA CS
- Int. Vac. Microelectron. Conf., 11th (1998), 32-33 Publisher: Institute of SO Electrical and Electronics Engineers, New York, N. Y. CODEN: 67IYAX
- DT Conference
- English LA
- UV photoelectron spectroscopic studies were made of Mo oxide, La Sr Co AΒ oxide, PZT, and Pb Nb Ti Zr oxide. The field emission was studied. The work functions are given.
- Dielectric films IT

Work function

(field emission and work function of oxide films on molybdenum emitter tips)

Oxides (inorganic), processes IT

RL: PEP (Physical, engineering or chemical process); PROC (Process) (field emission and work function of oxide films on molybdenum emitter tips)

- ANSWER 3 OF 11 HCAPLUS COPYRIGHT 2002 ACS T.49
- AN 1999:134892 HCAPLUS
- DN 130:275033
- Optoelectronic properties controlled by an electric field in thin films TI
- ΑU Olesik, J.; Calusinski, B.; Olesik, Z.
- Inst. Physics, Pedagogical Univ., Czestochowa, 42-201, Pol. CS
- Electron Technol. (1998), 31(3/4), 425-428 SO CODEN: ETNTAT; ISSN: 0070-9816
- PΒ Institute of Electron Technology
- DTJournal
- LA English
- Electron emission properties of semiconducting films (In203:Sn) AΒ and metallic films (Ti) of thickness 10-100 nm were studied. The films were deposited by reactive ion sputtering on a glass substrate of thickness 0.2 mm. The opposite side of the substrate (with a field electrode evapd. onto it) was biased by neg. voltage. This created transverse elec. field which favored electron emission into the vacuum. The study was performed in the vacuum of the order 10-8 hPa. Electron emission yield dependence on the intensity of an internal field and illumination were measured. The emission yield for semiconducting films depends exponentially on field intensity. The field influence on photoemission was also found. For metallic films the field effect on emission phenomena is significantly smaller. Measurements of electrons energy in field induced emission for both types of studied emitters showed that .apprx.80% of electrons have energy up to 10 eV but some electrons (a few percent) of energy .apprx.50 eV are also detected.
- Photoelectron spectrometers IT
 - (for measuring optoelectronic properties controlled by elec. field in thin films)
- Metals, properties TT
 - RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PROC (Process); USES (Uses)
- TΤ Ion beam sputtering Sputter deposition
 - (to evap. thin films of a semiconductor and metal

```
onto glass substrates)
    7440-32-6, Titanium, properties 50926-11-9, Indium tin oxide
    RL: PEP (Physical, engineering or chemical process); PRP (Properties); TEM
     (Technical or engineered material use); PROC (Process); USES (Uses)
        (optoelectronic properties controlled by elec. field in thin films of)
             THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 8
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L49 ANSWER 4 OF 11 HCAPLUS COPYRIGHT 2002 ACS
    1998:739386 HCAPLUS
AΝ
DN
    130:45059
ΤI
    Cold cathode fluorescent lamps
    Hirohashi, Masaki; Kuwata, Atsushi
IN
    Matsushita Electric Industrial Co., Ltd., Japan
PA
    Jpn. Kokai Tokkyo Koho, 7 pp.
     CODEN: JKXXAF
DT
    Patent
    Japanese
LΑ
FAN.CNT 1
                    KIND DATE
    PATENT NO.
                                          APPLICATION NO. DATE
     _____
                                          -----
                     A2
PΙ
    JP 10302714
                           19981113
                                          JP 1997-106989
                                                           19970424
    An electron-emitter coated on the cold cathode in the lamps
AB
    suitable for use in backlights comprises .gtoreq.1 oxygen-deficient
metal oxide, AlaA2bA3cBldB2eB3fB4gB5hOz, where Al = Ce;
     A2 = Y, La, Pr, Nd, Pm, Sm, Eu, Gd, Tb, Dy, Ho, Er, Tm, Yb, Lu; A3 = Ca,
     Sr, Ba; B1 = Cu; B2 = Cr, Mn, Fe, Co, Ni; B3 = Ti, Zr, Ru, Rh, Hf, Os, Ir;
     B4 = V, Nb, Ta; B5 = Mo, W; and 4a+3b+2c+2d+3e+4f+5g+6h > 2z.
     216577-19-4, Lanthanum manganese strontium oxide (La0.8Mn0.9Sr0.202.9)
ΙT
     216577-21-8, Calcium manganese yttrium oxide (Ca0.1MnY0.902.7)
     216577-22-9 216577-23-0, Ruthenium strontium yttrium oxide
     (Ru0.2SrY0.802.9)
     RL: DEV (Device component use); USES (Uses)
        (cold cathode fluorescent lamps)
L49 ANSWER 5 OF 11 HCAPLUS COPYRIGHT 2002 ACS
    1998:479005 HCAPLUS
ΑN
    129:128782
DN
    Selective infrared line emitters
ΤТ
    Chen, Zheng; Rose, Millard Franklin; Adair, Peter L.
ΤN
PA
    Auburn University, USA
    U.S., 7 pp.
SO
     CODEN: USXXAM
DT
     Patent
    English
LA
FAN.CNT 1
                    KIND DATE
     PATENT NO.
                                         APPLICATION NO. DATE
     US 5780370 A 19980714 US 1996-699509 19960819
    US 5780370 A
PΙ
     Selective IR line emitter comprising a non-woven composite of
     fibers of .gtoreq.1 rare earth metal oxide and
     .gtoreq.1 structure-forming material selected from the group consisting of
     alumina, silica, yttrium oxide, and zirconium oxide, where the
     rare earth metal oxide fibers are dispersed and
     interlocked in a network of the structure forming fibers, and where the
     fibers are adhesively connected at a multiplicity of crossing points by a
     ceramic bonding agent. The bonding agent may be a ceramic compd. formed
     by a sol-gel process. The articles exhibit narrow bandwidth emissions
     with good thermal conversion efficiencies and with improved tensile
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strength and lower brittleness than in the absence of the ceramic bonding agent. Application to heating systems is indicated. ΙT Ceramic composites IR sources Sol-gel processing (selective IR line emitters based on adhesively locked fiber Alumina fibers IT Ceramic fibers Silica fibers RL: DEV (Device component use); USES (Uses) (selective IR line emitters based on adhesively locked fiber IT Rare earth oxides RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (selective IR line emitters based on adhesively locked fiber composites) Heating systems IT (selective IR line emitters based on adhesively locked fiber composites for) 1303-86-2, Boria, uses TT 1313-97-9, Neodymium oxide. Zirconia, uses 1314-36-9, Yttria, uses 1314-37-0, Ytterbium oxide. 1344-28-1, Alumina, uses 7631-86-9, Silica, uses 12055-62-8, Holmium 12061-16-4, Erbium oxide. 37368-09-5, **Titanium** zirconium oxide RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (selective IR line emitters based on adhesively locked fiber composites) ANSWER 6 OF 11 HCAPLUS COPYRIGHT 2002 ACS L49 AN1996:294681 HCAPLUS DN 124:330095 TITransistors with oxide superconductor bases IN Toda, Norihiko; Abe, Hitoshi; Makita, Takehiko Oki Electric Ind Co Ltd, Japan PASO Jpn. Kokai Tokkyo Koho, 5 pp. CODEN: JKXXAF DTPatent Japanese LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ---------JP 08056022 A2 19960227 PΙ JP 1994-191614 19940815 Transistors contain insulator substrates (e.g., SrTiO3), semiconductor regions (contg. emitters) formed by implantation of metals (e.g., Nb), oxide superconductor base layers (e.g., (Ba, Rb)BiO3) partially covering the semiconductor regions, and collector regions (e.g., In) formed on the superconductor layers, and having wider area than the emitter regions. Carriers are effectively collected by the collectors. ΙT Superconductor devices (transistors with oxide superconductor bases and doped semiconductor emitters) IT Transistors (with oxide superconductor bases and doped semiconductor emitters) IT 122610-40-6, Barium bismuth rubidium oxide ((Ba,Rb)BiO3)

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RL: DEV (Device component use); USES (Uses)
        (transistors with oxide superconductor bases)
     7440-74-6, Indium, uses
    RL: DEV (Device component use); USES (Uses)
        (transistors with oxide superconductor bases and collectors from)
     7440-03-1, Niobium, uses
IT
     RL: MOA (Modifier or additive use); USES (Uses)
        (transistors with oxide superconductor bases and doped semiconductor
        emitters)
     12060-59-2, Strontium titanium oxide (SrTiO3)
IT
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (transistors with oxide superconductor bases on)
L49 ANSWER 7 OF 11 HCAPLUS COPYRIGHT 2002 ACS
    1995:573760 HCAPLUS
AN
    122:304660
DN
TТ
    Transistors with superconductor base
    Usuki, Tatsuro; Suzuki, Hiroshi; Yoshisato, Masanobu
TN
     Sanyo Electric Co, Japan
PA
     Jpn. Kokai Tokkyo Koho, 6 pp.
SO
     CODEN: JKXXAF
DT
     Patent
T.A
    Japanese
FAN.CNT 1
     PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
     _____ ____
PI
    JP 06283772
                      A2
                           19941007
                                          JP 1993-92261
                                                            19930325
     In fabrication of title transistor comprising a semiconductor collector
AΒ
     region, a superconductor base region formed on the collector region, and a
     emitter region formed on the base region via an insulating
     film using superconductor or metal, the thickness of the
     epitaxial insulating film in between the collector and the base is larger
     than 10 .mu.m, but less than the coherence length of the base region. The
     transparence of the quasi-particles is increased since the barrier in the
     interface of the base/collector is reduced due to the insulating film
     formed in this method.
IT
     Epitaxy
     Superconductor devices
     Transistors
        (fabrication of transistors with superconductor base)
TΤ
     1303-11-3, Indium arsenide, uses 1309-48-4, Magnesium monoxide, uses
     12795-57-2, Strontium titanium oxide
                                          107539-20-8,
     Yttrium barium copper oxide 118145-70-3, Barium bismuth potassium oxide
     RL: DEV (Device component use); USES (Uses)
        (fabrication of transistors with superconductor base)
    ANSWER 8 OF 11 HCAPLUS COPYRIGHT 2002 ACS
L49
     1995:561460 HCAPLUS
AN
     122:302406
DN
TI
    Light emitting device
     Takeda, Mikiro
IN
     Sharp Kk, Japan
PA
     Jpn. Kokai Tokkyo Koho, 5 pp.
SO
     CODEN: JKXXAF
DT
     Patent
LA
     Japanese
FAN.CNT 1
     PATENT NO.
                     KIND DATE
                                           APPLICATION NO. DATE
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_____
     _____
                    A2 19950131
                                         JP 1993-173341
                                                           19930713
    JP 07029682
PΤ
    The title device comprising a light emitting layer formed by a
AB
    semiconductor material, is characterized in that the light emitting
    layer is interposed between tunneling emitter
    layers as an accelerating region for carriers, formed by an
     insulator having a thickness less than the mean free path of carriers, and
    a ferroelec. layer is formed on the tunneling emitter as a
    currier source.
    Electroluminescent devices
IT
        (ferroelec. layer in electroluminescence device)
    12676-60-7, Lanthanum lead titanium zirconium oxide
IT
     (LaPbTiZrO3)
    RL: DEV (Device component use); USES (Uses)
        (ferroelec. layer in electroluminescence device)
     1314-98-3D, Zinc sulfide (ZnS), Mn activated
IT
    RL: DEV (Device component use); USES (Uses)
        (light emitting layer in electroluminescence device)
     12033-89-5, Silicon nitride (Si3N4), uses
IT
    RL: DEV (Device component use); USES (Uses)
        (tunneling emitter layer in
       electroluminescence device)
T.49
    ANSWER 9 OF 11 HCAPLUS COPYRIGHT 2002 ACS
AN
    1993:615790 HCAPLUS
DN
    119:215790
ΤI
    Low-voltage dielectric-base transistor
IN
    Tamura, Yasutaka
PA
    Fujitsu Ltd, Japan
    Jpn. Kokai Tokkyo Koho, 8 pp.
SO
    CODEN: JKXXAF
DT
    Patent
LA
    Japanese
FAN.CNT 1
    PATENT NO.
                     KIND DATE
                                         APPLICATION NO. DATE
                           19930806
     -----
                                          -----
    JP 05198855
                     A2
PΙ
                                         JP 1992-9028
                                                         19920122
    In the transistor comprising a high-dielec. const. base region, a base
AB
    electrode directly or indirectly contacting with the base region, and
     emitter and collector electrodes contacting with the base region
     through a barrier layer having a dielec. const. lower than that of the
    base region, the base, emitter, and/or collector electrode
    consists of the same substance as the base region, with the surface
     contacting with the barrier layer or the base region, which consists of an
     element-substituted or -deficient substance with elec. cond. The
     transistor may have a carrier tunneling barrier layer
    between the base region and the base electrode. The emitter or
     collector electrode may consist of a (super)conductive oxide. The base
     region may consist of an oxide contg. Sr, Ti, Ta, K, Sn, Zr, or Nb. The
    base region may consist of KTal-xNbxO3, and the base electrode may consist
    of KTal-xNbxO3 with K substituted for Ca. The transistor may comprise an
     insulating substrate covered with a thin film of the base region.
TT
    Transistors
        (dielec.-base, low-voltage)
IT
    Electric insulators and Dielectrics
        (low-voltage transistor base from)
     1306-38-3, Cerium dioxide, uses 12003-86-0, Aluminum yttrium oxide
IT
     (Alyo3)
    RL: USES (Uses)
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```
(low-voltage dielec.-base transistor carrier tunneling
       barrier layer)
     107539-20-8, Yttrium barium copper oxide
    RL: USES (Uses)
        (low-voltage dielec.-base transistor electrode)
     12060-59-2, Strontium titanium oxide (SrTiO3)
IT
     12710-39-3, Niobium potassium tantalum oxide
    RL: USES (Uses)
        (low-voltage transistor dielec. base and electrode from)
     7440-70-2, Calcium, uses
ΙT
    RL: USES (Uses)
        (potassium tantalate niobate doped with, for dielec.-base transistor
       base electrode)
TΤ
    7440-03-1, Niobium, uses
    RL: USES (Uses)
        (strontium titanate doped with, for dielec.-base transistor electrode)
    ANSWER 10 OF 11 HCAPLUS COPYRIGHT 2002 ACS
1.49
    1992:663052 HCAPLUS
AN
    117:263052
DN
    Oxide cathode emitter display tubes and manufacture of oxide
TI
     cathodes thereof
IN
    Kani, Akira; Sago, Sumuto; Iijima, Motoi; Yokoi, Tatsumasa; Kamiya,
    Hikonori; Asai, Hideyuki; Senda, Shinji; Kikuchi, Naoya; Matsuyama, Tatsuo
    Noritake Co., Ltd. K. K., Japan
PΑ
    Jpn. Kokai Tokkyo Koho, 8 pp.
SO
     CODEN: JKXXAF
DT
     Patent
    Japanese
LA
FAN.CNT 1
     PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
                     ----
                                           -----
                                          JP 1990-218947
PΙ
     JP 04104430
                      A2
                           19920406
                                                           19900822
                     B2
    JP 2525278
                           19960814
    The oxide cathode material useful in emitter display tubes is an
AΒ
     elec. conductive oxide of metals from Group IA, IIA,
     and/or IIIA elements. The manufg. the oxide cathode materials involves
     (1) pulverizing the conductive oxide, (2) kneeding a mixt. contg. the
    pulverized oxide, a liq. vehicle, and a binder to form a conductive paste,
     and (3) printing a cathode pattern with the paste on a substrate. The use
    of the oxides provides the cathode with a decreased operating voltage and
     an increased emission efficiency without Ag vapor as its sealing gas.
IT
    Cathode-ray tubes
        (elec. conductive oxide for)
    Electric conductors
IT
        (pastes, metal oxide for, for cathode
        emitter)
     12016-86-3, Lanthanum cobalt oxide (LaCoO3)
ΤT
                                                   12022-69-4, Strontium iron
     oxide (SrFeO3) 12031-18-4, Lanthanum nickel oxide (LaNiO3) 12031-41-3,
     Lanthanum nickel oxide (La2NiO4) 12035-28-8, Neodymium nickel oxide
               12047-25-5
                            12053-92-8, Copper lanthanum oxide (CuLa204)
     (Nd2NiO4)
     12143-36-1, Strontium vanadate (SrVO3)
                                             12201-04-6, Lanthanum titanate
              12313-89-2 37217-07-5, Lanthanum ruthenium oxide (La2Ru2O7)
     (LaTiO3)
     39282-77-4, Strontium chromate (SrCrO3) 107121-69-7, Cobalt lanthanum
                                     109466-64-0, Barium copper yttrium
     strontium oxide (CoLa0.7Sr0.3O3)
                         118819-36-6, Rubidium tungsten oxide (Rb0.1WO3)
     oxide (Ba2Cu3YO6.5)
     144815-63-4, Sodium titanium oxide (Na0.1TiO2)
     144856-75-7, Osmium scandium oxide (Os2Sc2O7)
    RL: TEM (Technical or engineered material use); USES (Uses)
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(conductive paste contg., for cathode in display tubes)

- L49 ANSWER 11 OF 11 HCAPLUS COPYRIGHT 2002 ACS
- AN 1967:14900 HCAPLUS
- DN 66:14900
- TI Tunnel emission from aluminum-aluminum **oxide metal** structures
- AU Hayashi, Takeshi; Nakano, Tomoyasu
- CS Matsushita Res. Inst. Tokyo Inc., Kawasaki, Japan
- SO Jpn. J. Appl. Phys. (1966), 5(10), 982-3 CODEN: JJAPA5
- DT Journal
- LA English
- Amethod of obtaining emission of electrons from cold materials by using a thin-film tunnel cathode in asym. diode structures is described. The diode consists of a 200-A. Al film (99.999%) on which an oxide layer is grown in O at 400.degree. for 2 hrs. A SiO film 1500 A. thick covers the edge of the active area of 1/160 cm.2 Surface emitter layers of Al 99.999%, Au 99.999%, Ti 99.9%, and Cr, 100-400 A. thick, are vacuum deposited on the active area. The current transfer is large with all the metals except Au. It is believed that all but the Au form a special oxide on the Al203 during the forming process, producing a heterojunction and increasing the tunnel effect.
- IT Diodes
 - (film, aluminum-aluminum oxide-metal,
 - tunnel emission in)
- IT Potential barriers
 - (tunnel emission through, from aluminum-aluminum oxidemetal structures)
- IT Electron emission
 - (tunnel, from aluminum-aluminum oxide-metal structures)
- IT Electric current
 - (tunnel, in aluminum-aluminum oxide-metal diodes)
- IT 7440-47-3, properties
 - RL: PRP (Properties)
 - (electron tunnel emission from aluminum-aluminum oxide-chromium structures)

```
L52 ANSWER 1 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    2002:69555 HCAPLUS
DN
    136:126657
    Field emission displays using diamond-like carbon thin layer
TI
    cathodes with high emission efficiency and good display images,
    and their manufacture
    Kim, Steven; Yom, Gun Yong; Lee, Do Hain
IN
PA
    Skion Corp., USA
    Jpn. Kokai Tokkyo Koho, 23 pp.
SO
    CODEN: JKXXAF
DТ
    Patent
    Japanese
LA
FAN.CNT 1
    PATENT NO.
                   KIND DATE
                                         APPLICATION NO. DATE
    ______
                                         -----
    JP 2002025426 A2 20020125
                                         JP 2001-112194
                                                        20010411
PΤ
US 2002011770
PRAI KR 2000-36066
                    A1 20020131
                                         US 2000-725164 20001129
                     Α
                          20000628
    The emitter comprises (A) a diamond-like carbon (DLC) thin
AB
    layer, (B) a lower electrode layer (maybe comprise Mo) having a
    1st hole tapered toward the DLC layer, (C) an insulator layer (maybe
    comprise SiO2) having a 2nd hole, which is larger than the 1st one and has
    a side wall rounded and tapered towards the electrode layer, above the 1st
    one, and (D) an upper electrode layer (maybe comprise Mo) having
    a 3rd hole smaller than the 1st one above the 1st one. The DLC layer may
    be deposited by Cs+ ion beam sputtering. An illustration of the emission
    device is given.
ΙT
    7439-98-7, Molybdenum, processes
    RL: CPS (Chemical process); DEV (Device component use); PEP (Physical,
    engineering or chemical process); PROC (Process); USES (Uses)
        (electrode layer; manuf. of field emission displays using diamond-like
       carbon thin layer cathodes with high emission
       efficiency and good display images)
    7782-44-7, Oxygen, uses 7782-50-5, Chlorine, uses
TT
    RL: NUU (Other use, unclassified); USES (Uses)
        (etching gas; manuf. of field emission displays using diamond-like
       carbon thin layer cathodes with high emission
       efficiency and good display images)
    ANSWER 2 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    2002:64135 HCAPLUS
AN
    136:94700
DN
    Manufacturing method of amorphous carbon field emitter with low
TI
    operating voltage
IN
    Lee, Nae Seong; Han, In Taek; Choi, Won Bong; Choi, Joon Hei
    Samsung Sdi Co., Ltd., S. Korea
PΑ
    Repub. Korean Kongkae Taeho Kongbo, No pp. given
SO
    CODEN: KRXXA7
DT
    Patent
    Korean
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                        APPLICATION NO. DATE
                   ----
                                         -----
    KR 2000009237 A
                          20000215
                                        KR 1998-29511
PΤ
                                                         19980722
    A manufg. method of a field emitter is provided to lower
    operating voltage required to electron emission and present a field
    emitter having an amorphous C micro-tip. The present invention
    discloses a manufg. method of a field emitter comprising: a step
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forming stripe shape cathode electrodes on substrate; a step forming an insulation layer on the cathode electrodes and the exposed substrate; a step forming a gate layer on the insulation layer; a step forming plural holes by etching from the top of the gate layer to the top of the cathode electrodes; a step forming a sepn. layer on the gate layer; a step forming micro-tips having prescribed height within the holes by using amorphous C particles; and a step removing the sepn. layer. Dielectric films

Etching

Field emission cathodes

Gate contacts

(manufg. method of amorphous carbon field emitter with low operating voltage)

IT 7440-44-0, Carbon, processes

RL: PEP (Physical, engineering or chemical process); PYP (Physical process); TEM (Technical or engineered material use); PROC (Process); USES

(manufg. method of amorphous carbon field emitter with low operating voltage)

ANSWER 3 OF 37 HCAPLUS COPYRIGHT 2002 ACS L52

2002:27644 HCAPLUS AN

136:94708 DN

- TTCold electron-emitting devices with lowered leakage current and method for their manufacture
- Hori, Junichi; Hiranaka, Koichi; Tottori, Satoru IN
- Matsushita Electric Industrial Co., Ltd., Japan PA
- Jpn. Kokai Tokkyo Koho, 8 pp. SO

CODEN: JKXXAF

DT Patent

Japanese LA

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PΙ JP 2002008518 A2 20020111 JP 2000-189522 20000623

- AB The device has a semiconductor layer (A), an insulation layer (B), a field emission transistor (FET) contg. a source region, a gate region, and a drain region, .gtoreq.1 emitter(s) (C) formed by etching the drain region or a part of A in contact with the drain region, and a low-resistivity layer (D, having a const. potential and resistivity .gtoreq.2 orders higher than that of a channel of the FET) below the source region and the gate region. The method is characterized in that a semiconductor layer below emitter(s) and that below a FET are formed in the same process. The low-resistivity layer may contain Ti, Cr, Al, Mo, Ta, Ni, Cu, Ag, Pd, W, and/or Sn or may be formed by adding Group V or Group III element to the semiconductor layer by ion implantation, plasma ion doping, or vapor phase growth method.
- ITField effect transistors Field emission cathodes

(manuf. of cold electron-emitting devices contg. emitters and FET with lowered leakage current)

IT 7429-90-5, Aluminum, processes 7439-98-7, Molybdenum, 7440-02-0, Nickel, processes 7440-05-3, Palladium, processes processes 7440-25-7, Tantalum, processes 7440-22-4, Silver, processes 7440-32-6, Titanium, processes 7440-31-5, Tin, processes 7440-33-7, Tungsten, processes 7440-36-0, Antimony, processes 7440-38-2, Arsenic, 7440-42-8, Boron, processes 7440-47-3, Chromium, processes 7440-50-8, Copper, processes 7440-55-3, Gallium, processes processes

IT

7440-74-6, Indium, processes 7723-14-0, Phosphorus, processes
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)
 (low-resistivity layer; manuf. of cold electron-emitting devices contg.
 emitters and FET with lowered leakage current)
7440-21-3, Silicon, processes 7440-44-0, Carbon, processes
7440-56-4, Germanium, processes
RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)

(semiconductor layer; manuf. of cold electron-emitting devices contg.

L52 ANSWER 4 OF 37 HCAPLUS COPYRIGHT 2002 ACS

AN 2001:812429 HCAPLUS

DN 136:78072

TI Fabrication of electron **emitter** device using polycrystalline diamond film

emitters and FET with lowered leakage current)

- AU Hatta, Akimitsu; Sumitomo, Taku; Inomoto, Hideo; Hiraki, Akio
- CS Department of Electronic and Photonic Systems Engineering, Kochi University of Technology, Kochi, 782-8502, Japan
- New Diamond and Frontier Carbon Technology (2001), 11(5), 307-312 CODEN: NDFTFF; ISSN: 1344-9931
- PB Scientific Publishing Division of MYU K.K.
- DT Journal
- LA English
- AB An electron emitter device using a polycryst. diamond film has been developed. The device is composed of a cathode of a polycryst. diamond film, gate electrodes of aluminum, and an insulating layer of a diamond-like carbon film. The diamond film was deposited on an n-type Si wafer by the conventional microwave plasma chem. vapor deposition (CVD) method, the diamond-like carbon film was deposited by an electron cyclotron resonance (ECR) plasma CVD method, and the aluminum electrodes were deposited by an electron beam evaporator and patterned by photolithog. The diamond-like carbon layer on the cathode was removed by oxygen plasma etching. Electron emission from the cathode to the gate electrodes was obsd. at a gate voltage of more than 30 V and the emission current was 0.5 mA at 40 V.
- IT 7440-44-0, Carbon, uses
 - RL: DEV (Device component use); USES (Uses)
 (diamond-like; electron emitter device with gate electrodes
 of aluminum and insulating layer of diamond-like carbon film)
- IT 7429-90-5, Aluminum, uses
 - RL: DEV (Device component use); USES (Uses)
 (electron emitter device with gate electrodes of aluminum and insulating layer of diamond-like carbon film)
- IT 7782-40-3, Diamond, properties
 - RL: DEV (Device component use); PRP (Properties); USES (Uses) (fabrication of electron **emitter** device using polycryst. diamond film)
- RE.CNT 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L52 ANSWER 5 OF 37 HCAPLUS COPYRIGHT 2002 ACS
- AN 2001:710064 HCAPLUS
- DN 135:265824
- TI Diamond-based cathodes having electron supply layers and showing large electron emission quantity
- IN Nishibayashi, Yoshiki; Imai, Takahiro; Fujimori, Naoharu; Kitabatake,

Serial No.:09/846,127

Makoto; Watanabe, Akihiko; Yoshikawa, Masanori Fine Ceramics Center, Japan; Sumitomo Electric Industries, Ltd.; PA Matsushita Electric Industrial Co., Ltd. Jpn. Kokai Tokkyo Koho, 9 pp. CODEN: JKXXAF DT Patent Japanese LA FAN.CNT 1 APPLICATION NO. DATE KIND DATE PATENT NO. _____ -----JP 2001266736 A2 20010928 JP 2000-85092 20000324 PΙ The cathodes have diamond-made emitters and electron supply AB layers formed on/in the diamond surfaces. The electron supply layers may be doped with B in high concn. The electron supply layers may be graphite or electroconductive amorphous carbon. Ion implantation IT (boron doping by; diamond-based cathodes having electron supply layers and showing large electron emission quantity) IT Cathodes (diamond-based cathodes having electron supply layers and showing large electron emission quantity) IT 7440-44-0P, Carbon, processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PNU (Preparation, unclassified); PREP (Preparation); PROC (Process); USES (Uses) (amorphous, electroconductive, electron-supply layers; diamond-based cathodes having electron supply layers and showing large electron emission quantity) ANSWER 6 OF 37 HCAPLUS COPYRIGHT 2002 ACS L52 2001:489832 HCAPLUS AN DN 135:84429 TIField emission device and method for fabricating the same ΙN Choi, Jun-Hee; Cha, Seung-Nam; Lee, Hang-Woo PΑ S. Korea U.S. Pat. Appl. Publ., 12 pp. SO CODEN: USXXCO DTPatent English LA FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE -----------US 2001006325 A1 20010705 EP 1115134 A1 20010711 US 2001-754275 PΙ 20010105 EP 2001-300051 20010104 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO JP 2001216887 A2 20010810 JP 2001-315 20010105 PRAI KR 2000-361 20000105 Α Field emission devices are described which comprise a substrate; a cathode formed over the substrate; microtips having nanosized surface features formed on the cathode; a gate insulation layer formed over the substrate with wells in each of which a single microtip is located; a gate electrode, formed on the gate insulation layer, with gates aligned with the wells so that each of the microtips is exposed through a corresponding gate; a focus gate insulation layer having openings to each of which one or more gates correspond formed on the gate electrode; and a focus gate electrode with focus gates aligned with the openings of the focus gate

insulation layer formed on the focus gate insulation layer. Methods for fabricating the devices are also described which entail forming the

cathode, gate insulation layer with wells, and gate electrode on a substrate in sequence, and forming microtips on the cathode exposed by the wells; forming a focus gate insulation layer on the gate electrode to have a predetd. thickness with a polymer (e.g., a polyimide or photoresist) layer so that the wells having the microtips are filled; forming a focus gate electrode on the focus gate electrode; forming a predetd. photoresist pattern on the focus gate electrode; etching the focus gate electrode into a focus gate electrode pattern using the photoresist pattern as a etch mask; etching the focus gate insulation layer exposed trough the focus gate electrode pattern by plasma etching to open wells; etching the polymer layer within the wells of the gate insulation layer by plasma etching so that the polymer layer partially remains on the surface of the microtips; and etching the surface of the microtips by plasma etching using the polymer layer remaining on the microtips as an etch mask, and etching the polymer layer itself, to produce microtips with nanosized surface features. The plasma etching is preferably conducted using oxygen or an oxygen-contg. mixt.

IT Optical imaging devices

(field emission displays; field-emission devices with microtips with nanosized features and their fabrication)

IT Electronic device fabrication

Field emission cathodes

Field emitters

Photoresists

(field-emission devices with microtips with nanosized features and their fabrication)

L52 ANSWER 7 OF 37 HCAPLUS COPYRIGHT 2002 ACS

AN 2001:474158 HCAPLUS

DN 135:68674

TI Field emission displays of cathode arrays with carbon nanotube films as emitter and methods for their fabrication

IN Han, Si Wook; Kim, Sang Mun

PA LG Electronics Inc., S. Korea

SO U.S. Pat. Appl. Publ., 10 pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

1111. 01.1 1							
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE			
PI 'US 2001004979	A1	20010628	US 2000-733952	20001212			
PRAI KR 1999-57590	Α	19991214					

Field emission displays are described which comprise a cathode array including a cathode electrode formed on a substrate; insulating layers and carbon nanotube films for use as emitter electrodes formed alternately on the cathode electrode; and, a gate electrode formed on the insulating layer. Methods for fabrication of the field emission displays are discussed which entail forming a cathode electrode, an insulating layer, and a gate electrode on a substrate in succession; etching the gate electrode and the insulating layer into a cathode array pattern, to form an emitter region; forming a sacrificial layer on the gate electrode which is not etched; depositing a carbon nanotube film on the cathode electrode in the emitter region, to form an emitter; and, etching the sacrificial layer for removing the carbon nanotube formed on the sacrificial layer, to form a cathode array. Alternative fabrication methods in which the sacrificial layer is formed on the substrate before the etching are also described.

04/08/2002

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7439-98-7, Molybdenum, processes 7440-02-0, Nickel, processes
    7440-06-4, Platinum, processes 7440-22-4, Silver, processes
    7440-47-3, Chromium, processes
                                     7440-50-8, Copper, processes
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (gate electrode; field emission displays of cathode arrays with carbon
       nanotube films as emitter, and methods for their fabrication)
    7440-44-0, Carbon, processes
TΤ
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (nanotubes, film; field emission displays of cathode arrays with carbon
       nanotube films as emitter, and methods for their fabrication)
    ANSWER 8 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    2001:421246 HCAPLUS
AN
DN
    135:26979
    Field emission displays having patternwise activated flat diamond cathodes
ΥT
    and their manufacture
    Tomokage, Hajime; Iseri, Yoichi; Kim, Ho Do; Choi, Woon
IN
    Foundation for Scientific Technology Promotion, Japan; Mitsumi Electric
PA
    Co., Ltd
SO
    Jpn. Kokai Tokkyo Koho, 5 pp.
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
                    KIND DATE
    PATENT NO.
                                          APPLICATION NO. DATE
     _____
                                          ------
    JP 2001160366
                     A2
                                          JP 1999-343232
                           20010612
PΙ
                                                           19991202
    The displays comprise substrates, patternwise activated diamond
AB
    cathode layers, and phosphor-coated transparent anode
    layers above the cathodes. In the manufg. process, the
    cathode layers are activated and stabilized by
    application of high elec. field in vacuo and then patternwise applied with
    concd. field by ball anodes to form patterns. The patterns are then
    projected on the anodes upon voltag
    7440-57-5, Gold, uses
    RL: NUU (Other use, unclassified); USES (Uses)
        (ball anodes; manuf. of field emission displays with
       patternwise-activated flat diamond cathodes)
    7782-40-3, Diamond, processes
IT
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (polycryst., B-doped, cathode layers; manuf. of
        field emission displays with patternwise-activated flat diamond
       cathodes)
IT
    7440-21-3, Silicon, uses
    RL: DEV (Device component use); USES (Uses)
        (substrates; manuf. of field emission displays with
       patternwise-activated flat diamond cathodes)
    ANSWER 9 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    2001:397200 HCAPLUS
ΑN
DN
    134:375073
ΤI
    Cathode structure for planar emitter field emission displays
IN
    Russ, Benjamin E.
PΑ
    Sony Electronics Inc., USA
SO
    PCT Int. Appl., 15 pp.
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CODEN: PIXXD2
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DT Patent LA English

LA English FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 2001039236 A1 20010531 WO 2000-US31631 20001117

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,

AB A cathode structure for use in field emission display (FED) devices includes 4 layers. A 1st layer consists of conducting lines supported on an insulating substrate. A 2nd layer consists of thin nonconducting lines crossing the conducting lines. A 3rd layer consists of a thick layer of nonconducting material with holes centered between the thin nonconducting lines of the 2nd layer and extending over a portion of the thin nonconducting lines. A 4th layer consists of conducting lines contg. holes of the same dimension as and aligned with the holes in the 3rd layer exposing portions of the conducting lines of the 1st layer and of the nonconducting lines of the 2nd layer. Emissive material is deposited on the exposed portions of the conducting lines of the 1st layer to produce a cathode for an FED device. The 4-layer cathode structure improves emission characteristics such as c.d. and uniformity for planar edge emitters and surface emitters.

IT Glass, uses

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(spin on, non-conductive line; cathode structure for planar emitter field emission displays)

IT 7429-90-5, Aluminum, uses 7440-47-3, **Chromium**, uses 7440-57-5, **Gold**, uses 11118-57-3, **Chromium** oxide 50926-11-9, Indium tin oxide

RL: DEV (Device component use); TEM (Technical or engineered material use); USES (Uses)

(conductive line; cathode structure for planar emitter field emission displays)

L52 ANSWER 10 OF 37 HCAPLUS COPYRIGHT 2002 ACS

AN 2001:320245 HCAPLUS

DN 134:335437

TI Method of fabricating a field emission device with a lateral thin-film edge emitter

IN Karpov, Leonid D.; Eaton, Mark F.

PA Stellar Display Corp., USA

SO PCT Int. Appl., 45 pp. CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI WO 2001031671 Al 20010503 WO 2000-US29584 20001026 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,

AB A novel edge emitter device including an anode and a cathode situated at a level above and laterally displaced from the anode, providing an opening for a window above the anode. The cathode has an emitting edge which is operable to emit field electrons when a pos. voltage is applied to the anode with respect to the cathode. On the top

IT

IT

TТ

IT

IT

IT

IT

IT

IT

IT

surface of the anode is disposed either a phosphor layer operable to luminesce when struck with the electrons emitted from the emitting edge or a layer having a higher secondary emission ratio than anode. The device is capable of being configured as a diode, triode, tetrode, etc. having .qtoreq.1 control electrodes to control the current from emitting edge to anode. A fabrication process is capable of automatic alignment of cathode above insulating layer and around the window above anode and for the protrusion of cathode slightly beyond insulating layer into the window opening. Nanotubes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (carbon, cathode layer; method of fabricating a 12070-14-3, Zirconium carbide RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (Cr-ZrC-Cr cathode layer; method of fabricating a field emission device with a lateral thin-film edge emitter) 7429-90-5, Aluminum, processes 7440-47-3, Chrome, processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (anode layer; method of fabricating a field emission device with a lateral thin-film edge **emitter**) 7782-40-3, Diamond, processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (cathode layer; method of fabricating a field emission device with a lateral thin-film edge emitter) 7440-02-0, Nickel, processes 11105-45-6 RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (conducting layer; method of fabricating a field emission device with a lateral thin-film edge **emitter**) 1344-28-1, Alumina, processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (insulating layer, secondary electron emission materials; method of fabricating a field emission device with a lateral thin-film edge emitter) 7631-86-9, Silica, processes 113443-18-8, Silicon oxide (SiO) RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (insulating layer; method of fabricating a field emission device with a lateral thin-film edge emitter) 7440-44-0, Carbon, processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (metal-carbon-metal cathode layer; method of fabricating a field emission device with a lateral thin-film edge emitter) 7440-50-8, Copper, processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (pillar with chrome cap; method of fabricating a field emission device with a lateral thin-film edge emitter) 7440-21-3, Silicon, processes

RL: DEV (Device component use); PEP (Physical, engineering or chemical

process); PROC (Process); USES (Uses)

IT

(resistive layer, amorphous; method of fabricating a field emission
 device with a lateral thin-film edge emitter)
409-21-2, Silicon carbide, processes 12033-62-4, Tantalum
nitride 25583-20-4, Titanium nitride

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(resistive layer; method of fabricating a field emission device with a lateral thin-film edge **emitter**)

1T 1304-28-5, Barium oxide, processes 1309-48-4, Magnesium oxide, processes
RL: DEV (Device component use); PEP (Physical, engineering or chemical
process); PROC (Process); USES (Uses)

(secondary electron emission materials; method of fabricating a field emission device with a lateral thin-film edge emitter)

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L52 ANSWER 11 OF 37 HCAPLUS COPYRIGHT 2002 ACS

AN 2000:887848 HCAPLUS

DN 134:65126

TI Manufacture of electron emission components and display devices

IN Shimamura, Toshinori; Inoue, Hiroshi; Tatezono, Shinichi; Yamagishi, Takeshi

PA Sony Corp., Japan; Hitachi Funmatsu Yakin Co., Ltd.

SO Jpn. Kokai Tokkyo Koho, 7 pp. CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI JP 2000353466 A2 20001219 JP 1999-162815 19990609

AB Carbon coatings contg. C particles and thermosetting resins are formed on resist films and cathode layers, IR radiation is applied from light-transmitting substrate side to the coating side, where gate electrode layers function as masks, enhancing the adhesion of the coatings and substrates, as well as that of the C particles and the resins, and the resist films are dissolved, forming

IT Cathodes

Coating process

Electrooptical imaging devices

crown-like emitters in openings.

(carbon coating in manuf. of electron emission components and display devices)

IT Particles

(carbon particles in manuf. of electron emission components and display devices)

IT IR radiation

(manuf. of electron emission components and display devices)

IT 7440-44-0, Carbon, processes

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(carbon coating in manuf. of electron emission components and display devices)

- L52 ANSWER 12 OF 37 HCAPLUS COPYRIGHT 2002 ACS
- AN 2000:877083 HCAPLUS
- DN 134:35150
- TI Electron emitters, manufacture of the emitters, and display having the emitters

```
04/08/2002
    Neqishi, Eisuke; Nakata, Satoshi
TN
    Sony Corp., Japan
PΑ
    Jpn. Kokai Tokkyo Koho, 9 pp.
SO
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
                    KIND DATE
                                         APPLICATION NO. DATE
    PATENT NO.
     -----
                                          -----
    JP 2000348601 A2 20001215
                                         JP 1999-159485 19990607
PΙ
    The electron emitters are characterized by that they can be
AΒ
     driven by low elec. field and that they are suitable for ultrathin film
     display. Fine holes, penetrating through an upper gate layer and an
     intermediate insulating layer and part of a lower cathode
     layer, are formed. The each hole has the bottom with bumpy
     texture, and an electron-emitting layer (e.g., a diamond-like carbon or
     amorphous carbon layer, preferably formed by filtered cathodic vacuum arc
     deposition, sputtering, or laser ablation) is formed on the bottom. The
     electron-emitting layer has the surface with bumpy texture because of the
    bottom of the hole. Many electrons can be emitted from the
     electron-emitting layer by a low elec. filed of .ltoreq.50 V/.mu.m,
    because the surface of the electron-emitting layer is lower than the
     surface of the cathode layer and the texture of the
     electron-emitting surface is bumpy. The electron-emitting layer is formed
     after formation of the intermediate insulating layer, so that the electron
     emitting layer is prevented from being damaged by the plasma in formation
     of the intermediate insulating layer.
IT
     Cathodes
     Laser ablation
     Sputtering
        (formation of electron emitter having surface with bumpy
        texture made of carbon formed by sputtering or laser ablation)
IT
     Vapor deposition process
        (plasma, vacuum, filtered cathodic; formation of electron
        emitter having surface with bumpy texture made of carbon formed
       by sputtering or laser ablation)
IT
    Electroluminescent devices
        (ultrathin film; formation of electron emitter having surface
       with bumpy texture made of carbon formed by sputtering or laser
        ablation for)
     7440-44-0, Carbon, processes
IT
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (formation of electron emitter having surface with bumpy
        texture made of carbon formed by sputtering or laser ablation)
L52 ANSWER 13 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    2000:861258 HCAPLUS
AN
    134:36041
    Manufacture of electron emission source for fluorescent display device
TI
    Ito, Shigeo; Yamaura, Tatsuo
IN
```

DN

PΑ Futaba Denshi Kogyo Co., Ltd., Japan

Jpn. Kokai Tokkyo Koho, 11 pp. SO CODEN: JKXXAF

DTPatent

Japanese LA

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE ------ ---- ----

```
04/08/2002
                           20001208
                                           JP 1999-145559
                     A2
                                                            19990525
     JP 2000340100
     The invention relates to a process for making an electron emission source
     for low-voltage operation, suitable for use in a fluorescent display
     device, wherein the process includes formation of carbon layer
     on a cathode layer from a carbon material selected
     from a group consisting of carbon nano-tube, fullerene, nanoparticle and
     nanocapsule.
TT
     Nanotubes
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (carbon; manuf. of electron emission source having carbon layer formed
        from specified carbon material for fluorescent display device)
TT
     Optical imaging devices
        (fluorescent; manuf. of electron emission source having carbon layer
        formed from specified carbon material for fluorescent display device)
IT
     Carbon fibers, processes
     Glass fibers, processes
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (manuf. of electron emission source for fluorescent display device
        contq.)
IT
     Field emission cathodes
     Nanoparticles
        (manuf. of electron emission source having carbon layer formed from
        specified carbon material for fluorescent display device)
IT
     Fullerenes
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (manuf. of electron emission source having carbon layer formed from
        specified carbon material for fluorescent display device)
     7440-44-0, Carbon, processes
                                  99685-96-8, Fullerene
IT
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (manuf. of electron emission source having carbon layer formed from
        specified carbon material for fluorescent display device)
TT
     7429-90-5, ALuminum, processes 7440-21-3, Silicon, processes
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
```

process); PROC (Process); USES (Uses)

(porous material; manuf. of electron emission source for fluorescent display device contg.)

- ANSWER 14 OF 37 HCAPLUS COPYRIGHT 2002 ACS T₁5.2
- AΝ 2000:804072 HCAPLUS
- DN 133:343594
- Emitter in field-emission cathode and manufacture of the TI
- Inoue, Kazunori; Betsui, Keiichi; Nakaya, Tadashi IN
- PΑ Fujitsu Ltd., Japan
- Jpn. Kokai Tokkyo Koho, 11 pp. SO CODEN: JKXXAF
- DT Patent
- T₁A Japanese
- FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE ----

- A2 20001114 PТ JP 2000315453 JP 1999-124461 19990430
- The emitter consists of a substrate and an isotropic carbon fine AB powder-based elec. conductive cathode fixed on the substrate. The emitter is manufd. by forming an adhesive layer on the substrate,

IT

TT

IT

IT

IT

IT

IT

IT

AN DN

ΤI

IN

PA

SO

DT

LA

PΤ

applying the isotropic carbon fine powder on the adhesive layer, firing the composite, and pattering the adhesive layer and the cathode material layer. Alternatively, a mixt. contg. the carbon powder and elec. conductive adhesive solvent is applied on the substrate, dried, fired, and patterned to give the emitter. Other methods using low m.p. glass or using transfer technique are also claimed. The emitter is suitable for large area field-emitting display. Adhesives (conductive; for formation of emitter in field-emission cathode made of isotropic powder) Solvents (elec. conductive adhesive; for formation of emitter in field-emission cathode made of isotropic powder) Adhesives (for formation of emitter in field-emission cathode made of isotropic powder) Field emission cathodes (formation of emitter in field-emission cathode made of isotropic powder) Etching Photolithography Transfers (in formation of emitter in field-emission cathode made of isotropic powder) Glass, processes RL: PEP (Physical, engineering or chemical process); PROC (Process) (low-m.p.; for formation of emitter in field-emission cathode made of isotropic powder) Solvents (org., volatile; for formation of emitter in field-emission cathode made of isotropic powder) 7440-44-0, Carbon, processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (formation of emitter in field-emission cathode made of isotropic powder) L52 ANSWER 15 OF 37 HCAPLUS COPYRIGHT 2002 ACS 2000:781126 HCAPLUS 133:357320 Manufacture of electron emitter used for luminescence optical imaging display device Yamaura, Tatsuo; Ito, Shigeo Futaba Denshi Kogyo Co., Ltd., Japan Jpn. Kokai Tokkyo Koho, 6 pp. CODEN: JKXXAF Patent Japanese FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE ____ JP 2000311590 A2 20001107 JP 1999-119071 19990427 The process comprises depositing a cathode conductor material on an insulating substrate, forming a carbon layer on the cathode conductor layer by applying a paste material contq. carbon nanotubes, a fullerene, nanoparticles, and/or nanocapsules, and depositing particles on the carbon layer, and removing the particles.

The carbon nanotubes are formed almost parallel to the substrate surface,

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thereby reducing the concn. of elec. field.
IT
    Cathodes
     Optical imaging devices
     Transfer printing
        (manuf. of electron emitter used for luminescence optical
        imaging display device)
IT
     Fullerenes
     RL: DEV (Device component use); USES (Uses)
        (manuf. of electron emitter used for luminescence optical
        imaging display device)
     7440-44-0, Carbon, uses
IT
     RL: DEV (Device component use); USES (Uses)
        (manuf. of electron emitter used for luminescence optical
        imaging display device)
    ANSWER 16 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    2000:420544 HCAPLUS
AN
DN
    133:67446
    Manufacture of field-emission cold cathodes
ΤI
     Seko, Nobuya
TN
    NEC Corp., Japan
PΑ
SO
     Jpn. Kokai Tokkyo Koho, 20 pp.
     CODEN: JKXXAF
DT
     Patent
LA
    Japanese
FAN.CNT 1
                     KIND DATE
     PATENT NO.
                                           APPLICATION NO. DATE
                     ----
                            -----
                                                           19981209
     JP 2000173448
                      A2
                           20000623
                                           JP 1998-349216
PΙ
     Double insulator layers are deposited on Si substrates, photoresists with
     openings are formed on the insulator layers, the both insulator layers are
     etched with the photoresists as masks to form openings, emitter
     materials (e.g, Mo) are deposited vertically, sacrificial layer
     materials (e.g., Al2O3) are vapor deposited from inclined direction while
     spinning the substrates, 2nd deposition layers are formed on the
     sacrificial layers, creating emitter top layer, and the
     sacrificial layers are etched to lift-off the 2nd deposition layers.
     cold cathodes thus manufd. have precision gate diam. smaller than resoln.
     limit of photolithog.
     Photolithography
IT
        (manuf. of field-emission cold cathodes)
IT
     Etching
        (of sacrificial layers in manuf. of field-emission cold cathodes)
     Field emission
IT
     Field emission cathodes
        (sacrificial layers in manuf. of field-emission cold
        cathodes)
     7439-98-7, Molybdenum, processes
IT
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (Mo deposition in manuf. of field-emission cold cathodes)
     7429-90-5, Aluminum, processes
TT
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PROC (Process); USES (Uses)
        (sacrificial layers in manuf. of field-emission cold cathodes)
    ANSWER 17 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
AN
     1999:779284 HCAPLUS
     132:8216
DN
```

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04/08/2002
    Field emission device using resistors and fabrication of same
ΤI
    Kim, Jong-Min
IN
    Samsung Display Devices Co., Ltd., S. Korea
PΑ
SO
    U.S., 11 pp.
    CODEN: USXXAM
DT
    Patent
    English
LA
FAN.CNT 1
                   KIND DATE
                                        APPLICATION NO. DATE
    PATENT NO.
    _____
                                         -----
    US 5998916
                    A 19991207
                                        US 1998-38050 19980311
ΡI
    The title field emission device comprises a substrate, cathodes formed on
ΑB
    the substrate, a resistor layer continuously formed on the cathodes,
    microtips formed on the resistor layer, an insulating layer formed on the
    resistor layer and the substrate, and a gate formed on the insulating
    layer, wherein the resistor layer is formed by depositing diamond-like
    carbon (DLC) on the cathodes with PECVD. Accordingly, fabrication yield
    is high since the adhesion of the resistor layer to the
    cathodes is improved. Various types of resistor layers can be
    formed since the resistor layer has excellent chem. durability.
    reliability and consistency of the fabrication process is improved since
    the doping level is easily controlled.
ΙT
    Field emission
    Field emission cathodes
    Field emitters
    Resistors
       (field emission device using resistors and fabrication of same)
ΙT
    Vapor deposition process
       (plasma; field emission device using resistors and fabrication of same)
    7440-44-0, Carbon, processes
ΙT
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
       (diamond-like; field emission device using resistors and fabrication of
       same)
             THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 1
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L52 ANSWER 18 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    1998:735109 HCAPLUS
AN
    130:9727
DN
    Field-emission cathode
TI
    Itoh, Shigeo; Watanabe, Teruo; Ohtsu, Kazuyoshi; Taniguchi, Masateru
ΙN
PA
    Futaba Denshi Kogyo K.K., Japan
    U.S., 9 pp.
SO
    CODEN: USXXAM
DT
    Patent
LΑ
    English
FAN.CNT 1
                   KIND DATE
    PATENT NO.
                                        APPLICATION NO. DATE
     -----
                                         -----
                    Α
                                        US 1996-761134
PΤ
    US 5834885
                          19981110
                                                        19961206
    A field emission cathode which is capable of increasing bond strength
AB
    between emitters and a resistive layer includes a laminated
    board, which includes a substrate, and at least a cathode
    layer, a resistive layer, an insulating layer, and a gate
```

electrode layer which are deposited in the form of films on the substrate in order. The gate electrode layer and insulating layer are formed with through holes. The cathode also includes buffer layers made of an insulating material and formed on portions of the resistive layer exposed

```
via the through holes, as well as emitters arranged on the
    buffer layers, resulting in increased bond strength between the resistive
     layer and the emitters.
IT
    Buffers
    Electric conductors
    Electric insulators
     Resistors
     Semiconductor materials
        (field emission cathodes contg.)
TΤ
    Carbides
    Nitrides
    Refractory metals
    RL: DEV (Device component use); USES (Uses)
        (field emission cathodes contg.)
     Field emission cathodes
IT
        (having increased bond strength between emitters and
        resistive layers)
     7440-21-3, Silicon, uses
IT
     RL: DEV (Device component use); USES (Uses)
        (amorphous; field emission cathodes contg.)
     7440-03-1, Niobium, uses 7440-44-0, Carbon, uses
                                                        7631-86-9,
IT
     Silica, uses
     RL: DEV (Device component use); USES (Uses)
        (field emission cathodes contg.)
RE.CNT
             THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L52 ANSWER 19 OF 37 HCAPLUS COPYRIGHT 2002 ACS
     1998:334089 HCAPLUS
AN
DN
    129:61740
    Manufacture of field electron emitters
ΤI
     Kimura, Hideyoshi; Kondo, Yukihiro; Nishioka, Koji
IN
PA
    Matsushita Electric Works, Ltd., Japan
SO
     Jpn. Kokai Tokkyo Koho, 5 pp.
     CODEN: JKXXAF
DT
     Patent
    Japanese
LA
FAN.CNT 1
                     KIND DATE
     PATENT NO.
                                          APPLICATION NO. DATE
     ----- ----
                           ------
                                           -----
                                           JP 1996-286408
     JP 10134703
                      A2
                           19980522
PI
                                                            19961029
     Insulator layers surrounding cathode layers
AΒ
     have multilayer structures using materials (e.g., SiO2 and Al2O3) of
     different etching rates and having uneven sidewalls. Electron-emission-
     improving films (e.g., Cr) are vapor deposited on cathodes.
IT
     Cathodes
     Electric field
     Electron emission
        (manuf. of field electron emitters with multilayer insulator
        sidewalls)
     Dielectric films
ΙT
     Vapor deposition process
        (manuf. of field electron emitters with multilayer insulator
        sidewalls and vapor-deposited emission-improving films)
     7440-47-3, Chromium, uses
TT
     RL: DEV (Device component use); USES (Uses)
        (manuf. of field electron emitters with multilayer insulator
        sidewalls and emission-improving films from)
TT
     1344-28-1, Alumina, uses 7631-86-9, Silica, uses
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RL: DEV (Device component use); USES (Uses)

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(manuf. of field electron emitters with multilayer insulator
       sidewalls and vapor-deposited emission-improving films)
    ANSWER 20 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    1998:201892 HCAPLUS
AN
    128:315938
DN
    Fabrication and simulation of a gated thin film emitter
ΤI
ΑU
    Ko, Tae-Young; Chung, Bokeon; Lee, J. Y.; Jeon, D.
    Department of Physics, Myong Ji University, Yongin Kyunggi-Do, Seoul,
CS
    449-728, S. Korea
    J. Vac. Sci. Technol., B (1998), 16(2), 700-704
SO
    CODEN: JVTBD9; ISSN: 0734-211X
PΒ
    American Institute of Physics
    Journal
DT
    English
LΑ
    The authors have fabricated a gated field emitter using a
AB
    diamond-like carbon (DLC) film cathode. The process involved the
    deposition of DLC, insulator, and gate layers followed by back etching to
    expose a patterned DLC. The authors also simulated the emission behavior
    of the gated DLC cathode. The emission sites on the DLC film were
    simulated by multiple sharp points formed on the DLC surface. The
    electron trajectory and the emission current were studied as a function of
    structural parameters such as cathode height, oxide
    layer thickness, gate hole diam., and focus electrode.
    Physicochemical simulation
IT
        (fabrication and simulation of gated thin film emitter)
ΙT
    Field emission cathodes
        (thin-film; fabrication and simulation of gated thin film
       emitter)
    7440-44-0, Carbon, uses
IT
    RL: DEV (Device component use); USES (Uses)
        (diamond-like; fabrication and simulation of gated thin film
       emitter)
    7782-40-3, Diamond, uses
IT
    RL: DEV (Device component use); USES (Uses)
        (field emitter; fabrication and simulation of gated thin film
       emitter)
    ANSWER 21 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    1997:717545 HCAPLUS
AN
DN
    128:9519
ΤI
    Electrical field emitters
    Ito, Shigeo; Nakada, Hisashi; Iwasa, Tadashi
ΙN
    Futaba Denshi Kogyo Co., Ltd., Japan
PA
    Jpn. Kokai Tokkyo Koho, 9 pp.
SO
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                         APPLICATION NO. DATE
    ------ ---- -----
                                          -----
    JP 09283007
                     A2
                           19971031
                                          JP 1996-111151
PΤ
                                                           19960409
    JP 3235461
                     B2 20011204
AΒ
    The title emitters comprise a glass cathode substrate, a
    cathode electrode layer formed on the substrate, a pl.
    no. of conical emitters each formed apart on the cathode
    electrode, an insulator layer between the emitters on the
```

cathode electrode, and 1st gate electrodes formed on the insulator layer

to give a field emission cathode unit. The **emitters** further comprise a perforated 2nd gate electrode as a cutoff electrode provided above the cathode unit and an anode electrode provided further above. The same voltage is impressed on the 1st and 2nd gate electrodes to give a space charge satn. region so as to attract electrons from the region to the anode electrode by impressing a few tens of V on the anode electrode. The arrangement consequently gives the **emitters** a significantly low anode voltage.

IT Electrodes

(cutoff; elec. field emitters)

IT Field emission

Field emission cathodes

(elec. field emitters)

IT Space charge

(satn.; elec. field emitters)

IT 7429-90-5, Aluminum, properties

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process); USES (Uses)

(cathode; elec. field emitters)

IT 7439-98-7, Molybdenum, properties

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process); USES (Uses)

(conical emitter; elec. field emitters)

IT 50926-11-9, ITO

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process); USES (Uses) (transparent anode; elec. field emitters)

- L52 ANSWER 22 OF 37 HCAPLUS COPYRIGHT 2002 ACS
- AN 1997:275548 HCAPLUS
- DN 127:27075
- TI Fabrication of metal field **emitter** arrays on polycrystalline silicon
- AU Kim, II Hwan; Lee, Chun Gyoo; Kim, Yeo Hwan; Park, Byung Gook; Lee, Jong Duk
- CS Inter-University Semiconductor Research Center (ISRC) and School of Electrical Engineering, Seoul National University, Seoul, 151-742, S. Korea
- SO J. Vac. Sci. Technol., B (1997), 15(2), 468-471 CODEN: JVTBD9; ISSN: 0734-211X
- PB American Institute of Physics
- DT Journal
- LA English
- Cone-shaped metal field **emitter** arrays were fabricated on a single heavily doped polysilicon layer. From the observation of the metal field **emitter** fabricated on the single-polysilicon layer, it was noticed that the irregular shape of the gate aperture was caused by the large grain size of heavily doped polysilicon after oxidn. Therefore, in order to obtain a good shape, the double layers of polysilicon where one was heavily doped and the other was not doped were used. The undoped polysilicon layer was consumed for the gate oxide layer and the doped polysilicon layer was used to serve as a conducting **cathode**layer. The small grain size of the undoped polysilicon after oxidn. led to an improved shape of the gate aperture which rendered a stable emission characteristic.
- IT Field emission cathodes

(fabrication of metal field **emitter** arrays on polycryst. silicon)

IT 7723-14-0, Phosphorus, uses

```
RL: MOA (Modifier or additive use); USES (Uses)
        (doping; fabrication of metal field emitter arrays on
       polycryst. silicon)
    7439-98-7, Molybdenum, properties
                                       7440-21-3, Silicon,
IT
    properties
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PRP (Properties); PROC (Process); USES (Uses)
        (fabrication of metal field emitter arrays on polycryst.
       silicon)
L52 ANSWER 23 OF 37 HCAPLUS COPYRIGHT 2002 ACS
ΑN
    1996:598918 HCAPLUS
DN
    125:236040
    Manufacture of vertical fine cold cathodes
TI
    Yoshiki, Masayuki
IN
    Nippon Electric Co, Japan
PΑ
SO
    Jpn. Kokai Tokkyo Koho, 3 pp.
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                          APPLICATION NO. DATE
     -----
                           -----
                                          -----
    JP 08185795
                     A2
                           19960716
                                          JP 1994-324561 19941227
PΙ
                          19980408
                     B2
    JP 2737675
    The title process comprises formation of a protruding step on a Si
AB
    substrate, sequential lamination of a 1st insulating, a 1st gate
    conductive, a 2nd insulating, an emitter conductive, a 3rd
     insulating, and a 2nd gate conductive layer on the substrate, selective
    removal of the laminate on the step leaving the layers on the sides of the
    step and exposing the cross sections of the layers, and etching of the 2nd
    or the 3rd insulating layer deeper than the emitter conductive
     layer and the emitter conductive layer making the ends of the
    layer sharp.
    Cathodes
TΤ
        (field-emission, thin emitter layers with sharpened tips for
       silicon vertical field emitters)
    7439-98-7, Molybdenum, processes
IT
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (film; for emitter layers of vertical fine cold cathodes)
IT
    12627-41-7, Tungsten silicide
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (film; for gate layers of vertical fine cold cathodes)
IT
     7440-21-3, Silicon, uses
    RL: DEV (Device component use); USES (Uses)
        (thin emitter layers with sharpened tips for silicon vertical
        field emitters)
    ANSWER 24 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    1996:527611 HCAPLUS
AN
ĎΝ
    125:156379
TI
    Changing the surface quality of field emitters
    Ochiai, Hisataka; Ito, Shigeo
IN
PΑ
    Futaba Denshi Kogyo Kk, Japan
SO
    Jpn. Kokai Tokkyo Koho, 5 pp.
    CODEN: JKXXAF
DΤ
    Patent
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Japanese
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                         APPLICATION NO. DATE
     _____
                                         -----
    JP 08148083
                    A2 19960607
                                         JP 1994-311343 19941122
                     B2 19990203
    JP 2852356
    In field-emission-type cathodes, in which cathode layers
AB
     , insulator films, and gate electrode layers are successively formed on
     substrates, holes are formed in the electrode and insulator layers, and
     emitters are formed in the holes, the holes are filled with
     resists, the resists on the emitters are removed, the exposed
     emitters are coated with materials (e.g., TiN) with small work
     function, and the remaining resists are peeled off.
TT
    Work function
        (coating to change the surface quality of field emitters by
       materials of small work function)
    Coating materials
TT
        (to change the surface quality of field emitters by materials
       of small work function)
IT
    Cathodes
        (field-emission, coating to change the surface quality of)
     7440-25-7, Tantalum, uses 7440-33-7, Tungsten, uses
IT
     RL: DEV (Device component use); USES (Uses)
     (coating to change the surface quality of field emitters) 25583-20-4, Titanium nitride
IT
     RL: DEV (Device component use); USES (Uses)
        (coating to change the surface quality of field emitters by
       materials of small work function)
L52 ANSWER 25 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    1996:444117 HCAPLUS
AN
DN
    125:102817
TI
    Electron emitting devices
    Tomihari, Yoshinori
IN
    Kansai Nippon Electric, Japan
PA
     Jpn. Kokai Tokkyo Koho, 4 pp.
SO
     CODEN: JKXXAF
DT
     Patent
    Japanese
LA
FAN.CNT 1
     PATENT NO.
                    KIND DATE
                                        APPLICATION NO. DATE
     -----
                                         _____
                    A2
PΤ
     JP 08129951
                           19960521
                                         JP 1994-265044
                                                          19941028
                     B2
     JP 2964885
                           19991018
                     Α
                                         US 1995-547879 19951025
    US 5814926
                           19980929
PRAI JP 1994-265044
                          19941028
     The devices contain conical cathodes formed on substrates, lower insulator
     layers circling the cathodes with cavities in between,
     gate electrodes on the insulator layers, upper insulator layers on the
     gate electrodes, and electron coverging electrodes on the top, part of
     which is eccentric regarding the cathodes.
IT
    Electron sources
        (electron emitting devices with focusing electrodes)
IT
     7439-98-7, Molybdenum, uses 7440-21-3, Silicon, uses
     7440-33-7, Tungsten, uses
     RL: DEV (Device component use); USES (Uses)
        (electron emitting devices with focusing electrodes)
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L52 ANSWER 26 OF 37 HCAPLUS COPYRIGHT 2002 ACS

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1996:237612 HCAPLUS
AN
    124:276250
DN
    Field emission cathodes and manufacture thereof
TI
    Nakatani, Tadashi; Betsui, Keiichi; Fukuda, Shinya; Toyoda, Osamu
IN
PΑ
    Fujitsu Ltd, Japan
    Jpn. Kokai Tokkyo Koho, 10 pp.
SO
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
                    KIND DATE
                                          APPLICATION NO. DATE
    PATENT NO.
     _____
                                          -----
    JP 08017331 A2 19960119 JP 1994-149910 19940630
PΙ
    The cathode has an insulating film and a gate electrode having an opening
AΒ
    on the substrate or an emitter electrode thereon, and a cathode
    film consisting of an lower and an upper layer with inclination of edges
    of the latter (e.g., from Mo, Si, Ti, W) larger than that of the
    former (e.g., from Ni, Au, or Pt). A high resistance
    layer may be placed between the upper and the lower cathode
    layer or between the lower cathode layer and
    the substrate or the emitter electrode thereon. The title
    process comprises inclined incidence evapn. deposition of a sacrifice
    layer on the gate electrode without closure of the opening, and sequential
    vertical inclination evapn. deposition of the lower cathode
    layer without closure of the opening and the upper cathode
    layer resulting in closure of the opening. The emitter
    tip is formed to a small radius of curvature and electrons can be taken
    out at a lower voltage.
IT
    Electron sources
        (Mo and Mo silicide film for gate electrodes for
       metallic field emission cathodes)
TT
    Cathodes
        (field-emission, Mo and Mo silicide film for gate
        electrodes for metallic field emission cathodes)
                                7440-21-3, Silicon, uses
     7440-06-4, Platinum, uses
     7440-32-6, Titanium, uses 7440-33-7, Tungsten, uses
                                                            7440-57-5,
    Gold, uses
    RL: DEV (Device component use); USES (Uses)
        (Mo and Mo silicide film for gate electrodes for
        metallic field emission cathodes)
    7439-98-7P, Molybdenum, processes
ΙT
                                        7440-02-0P, Nickel,
               11104-85-1P, Molybdenum silicide
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process);
    USES (Uses)
        (Mo and Mo silicide film for gate electrodes for
        metallic field emission cathodes)
    ANSWER 27 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    1995:818973 HCAPLUS
AN
    123:244612
DN
    Field-emission cathodes and their manufacture
ΤI
    Ito, Shigeo; Yamada, Akira
IN
     Futaba Denshi Kogyo Kk, Japan
PΑ
    Jpn. Kokai Tokkyo Koho, 8 pp.
SO
    CODEN: JKXXAF
DT
     Patent
LA
    Japanese
FAN.CNT 1
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PATENT NO.
                                         APPLICATION NO. DATE
                   KIND DATE
     _____
                                          ______
    JP 07176264
                    A2 19950714
                                          JP 1993-344480
                                                          19931220
ΡI
    JP 2734965
                     B2 19980402
    The field-emission cathodes comprise substrates, cathode
AB
    layers, insulating layers, gate electrodes, and
    emitters formed in etched opening in the insulators; the
    emitters are prepd. by (1) ion-beam evapn. of low-m.p. metals and
     (2) surface deposition of metals by electron-beam evapn. or sputtering to
    form a tip. The cathodes are manufd. at low cost.
    Metals, processes
IT
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (field-emission cathodes)
    Ion beams
IT
    Sputtering
       (in manuf. of field-emission cathodes)
    Evaporation
ΙT
        (electron-beam, in manuf. of field-emission cathodes)
    Cathodes
IT
       (field-emission, particle-beam evapn. and sputtering in formation of
       tips for)
TT
    7429-90-5, Aluminum, processes 7439-89-6, Iron, processes
                                                                 7439-91-0.
    Lanthanum, processes 7439-95-4, Magnesium, processes Manganese, processes 7439-98-7, Molybdenum, processes
                                                            7439-96-5,
    7440-02-0, Nickel, processes 7440-03-1, Niobium, processes
                                                                 7440-05-3.
    Palladium, processes 7440-06-4, Platinum, processes
    7440-15-5, Rhenium, processes 7440-16-6, Rhodium, processes
                                                                  7440-18-8,
    Ruthenium, processes 7440-25-7, Tantalum, processes
    7440-26-8, Technetium, processes 7440-29-1, Thorium, processes
    7440-31-5, Tin, processes 7440-32-6, Titanium, processes 7440-33-7,
    Tungsten, processes 7440-44-0, Carbon, processes 7440-47-3,
    Chromium, processes 7440-50-8, Copper, processes 7440-57-5,
    Gold, processes 7440-61-1, Uranium, processes 7440-62-2,
    Vanadium, processes
                        7440-66-6, Zinc, processes 7440-67-7, Zirconium,
    processes
    RL: DEV (Device component use); PEP (Physical, engineering or chemical
    process); PROC (Process); USES (Uses)
        (field-emission cathodes)
    ANSWER 28 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    1995:613159 HCAPLUS
AN
DN
    123:24173
TI
    Field-emission cathodes
IN
    Ito, Shigeo
PΑ
    Futaba Denshi Kogyo Kk, Japan
SO
    Jpn. Kokai Tokkyo Koho, 7 pp.
    CODEN: JKXXAF
DT
    Patent
LA
    Japanese
FAN.CNT 1
                    KIND DATE
    PATENT NO.
                                          APPLICATION NO. DATE
     -----
                                          -----
                     A2
                                          JP 1993-260389
PΙ
    JP 07094075
                           19950407
    The spindle-type field-emission cathodes (having corn-shaped
AΒ
     emitters on a cathode electrode through a resistance layer) have
     conducting insulators between the resistance layer and the
     emitters, which are insulated by short-circuit current between the
     gate and the emitters. The conducting insulators may be formed
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on each emitter. The conducting insulators comprising Al, Aq,
    MnO2, or (p-n junction) Se may be insulated through electromigration. The
    resistance layer may comprise amorphous Si or Ta205.
IT
        (field-emission, field-emission cathodes with conducting insulator on
     1313-13-9, Manganese dioxide, uses
                                         7429-90-5, Aluminum, uses
IT
     7440-22-4, Silver, uses 7782-49-2, Selenium, uses
     RL: DEV (Device component use); USES (Uses)
        (field-emission cathodes with conducting insulator on emitters
     1314-61-0, Tantalum oxide (Ta2O5)
                                        7440-21-3, Silicon, uses
ΙT
     RL: DEV (Device component use); USES (Uses)
        (resistance layers; field-emission cathodes with
       conducting insulator on emitters)
    ANSWER 29 OF 37 HCAPLUS COPYRIGHT 2002 ACS
L52
    1995:584484 HCAPLUS
AN
DN
    123:46158
    Cathode for gaseous electric discharge panel and formation of the cathode
ТT
    Fujii, Kozo; Sawai, Hideo; Koiwa, Ichiro; Terao, Yoshitaka; Yamagata,
TN
    Toshikazu
PΑ
    Oki Electric Ind Co Ltd, Japan
SO
    Jpn. Kokai Tokkyo Koho, 7 pp.
     CODEN: JKXXAF
DT
    Patent
LA
    Japanese
FAN.CNT 1
                     KIND DATE
                                          APPLICATION NO. DATE
    PATENT NO.
     _____ ___
    JP 07085799
                     A2
                           19950331
                                          JP 1993-227647
PΙ
                                                           19930914
    The cathode comprises a cathode mother material, elec. conductive oxide
     layer having multiple pores piercing from surface of the oxide layer to
     inner part of the oxide layer, and an electron emitter substance
     crystal of an alk. earth metal oxide impregnated in the pores. Formation
    of the cathode involves the following steps; (1) forming the elec.
     conductive oxide layer on the cathode mother material,
     (2) forming the multiple pores in the oxide layer, (3) impregnating the
     the electron emitter substance in the pores, and (4) crystq. the
     electron emitter substance. The cathode is capable of
     discharging by low operating voltage.
IΤ
    Cathodes
        (discharge, for gaseous discharge panel)
    Optical imaging devices
IT
        (elec.-discharge, cathodes for)
     120897-67-8P, Calcium chromium lanthanum oxide (CaCrLaO3)
IT
     164294-50-2P, Calcium chromium cobalt lanthanum oxide
     (CaCrCoLaO3)
     RL: DEV (Device component use); PNU (Preparation, unclassified); PREP
     (Preparation); USES (Uses)
        (cathode elec. conductive layer; cathodes
        for gaseous discharge panel)
     68338-92-1, Chromium 6, iron 52, nickel 42
IT
    RL: DEV (Device component use); USES (Uses)
        (cathode mother material; cathodes for gaseous discharge panel)
ŦΤ
     12004-04-5P, Barium aluminate (BaAl2O4)
     RL: DEV (Device component use); PNU (Preparation, unclassified); PREP
     (Preparation); USES (Uses)
       (electron emitter in cathode; cathodes for gaseous discharge
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panel)

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L52 ANSWER 30 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    1995:330517 HCAPLUS
ΑN
DN
    122:94649
    Manufacture of field emission cathodes
TI
    Ito, Shiqeo; Watanabe, Teruo; Ochiai, Hisataka; Ootsu, Kazuyoshi;
    Taniguchi, Masateru
PA
    Futaba Denshi Kogyo Kk, Japan
    Jpn. Kokai Tokkyo Koho, 5 pp.
so
    CODEN: JKXXAF
    Patent
DT
    Japanese
LA
FAN.CNT 1
                    KIND DATE
                                        APPLICATION NO. DATE
    PATENT NO.
    _____
                                         -----
                   A2
                         19940506
    JP 06124649
                                         JP 1992-270580 19921008
PΙ
    JP 3180466
                     B2 20010625
    Title field emission cathode is manufd. by forming cathode
AB
    conductor layer, insulator layer and gate layer on an insulating
    substrate, opening holes through the insulator and the gate layers
    , anodizing the cathode conductors in the holes to form
    resistance layers in an electrolyte in which the cathode
    conductor layer is used as the anode and a passive electrode as
    the cathode, then forming cone-shaped Mo emitters on
    the resistance layers. This simple manuf. process provide uniform
    independent resistance layer for each emitter.
ΙT
    Anodization
       (manuf. of field emission cathodes)
IT
    Cathodes
       (field-emission, manuf. of)
IT
    7439-98-7, Molybdenum, uses 7440-25-7, Tantalum,
    uses 7631-86-9, Silica, uses
    RL: DEV (Device component use); USES (Uses)
       (manuf. of field emission cathodes contg.)
    1314-61-0, Tantalum oxide (ta2o5)
IT
    RL: DEV (Device component use); TEM (Technical or engineered material
    use); USES (Uses)
       (manuf. of field emission cathodes contg.)
L52 ANSWER 31 OF 37 HCAPLUS COPYRIGHT 2002 ACS
AN
    1994:691018 HCAPLUS
DN
    121:291018
тT
    Luminescent cathode emitter tubes and manufacture of them
    Kashiwakura, Yasuhide; Kanehisa, Osamu; Morishita, Hajime; Matsukyo,
ΤN
    Hideji; Shiiki, Masatoshi; Toyama, Hisashi; Morita, Yasukazu; Nakayama,
    Tsunekichi
    Hitachi Ltd, Japan; Hitachi Device Eng
PA
    Jpn. Kokai Tokkyo Koho, 12 pp.
SO
    CODEN: JKXXAF
DT
    Patent
    Japanese
LA
FAN.CNT 1
    PATENT NO.
                    KIND DATE
                                        APPLICATION NO. DATE
    -----
                                         -----
                                                        19920918
PΙ
    JP 06103893
                    A2 19940415
                                         JP 1992-248397
    Title manufg. involves forming an adhesive film on the inside surface of
AB
    its face plate, depositing a fluorescent particle layer on the adhesive
    layer, and vibrating to give the particle layer an increased bulk d. and
```

adhesion, wherein the adhesive contains a deliquescent inorg. compd. and an aq.-sol. polymer. The manufg. gives the bulbs an increased fluorescent particle layer d. and decreased pinholes.

IT Luminescence, cathodo-

(bulb; fluorescent particle layer coating on)

IT Adhesives

(deliquescent inorg. compd. and aq.-sol. polymer. for fluorescent particle layer on cathode emitter tube)

IT Coating materials

(luminescent, for fluorescent particle layer on cathode emitter tube)

IT 584-08-7, Potassium carbonate 9002-89-5, Poly(vinyl alcohol) 9003-05-8, Poly(acrylamide) 9003-39-8, Poly(n-vinylpyrrolidone) 9005-32-7, Alginate 9005-37-2, Propylene glycol alginate 9011-16-9D, Methyl vinyl ether-maleic anhydride copolymer, alkyl monoester with 10279-63-7, Potassium chromium sulfate 25231-54-3, Acrylamide-diacetone acrylamide copolymer 25322-68-3, Polyethylene glycol

RL: DEV (Device component use); MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (adhesive mixt. for fluorescent particle coating on cathode emitter tubes)

IT 7447-41-8, Lithium chloride, processes 7646-85-7, Zinc chloride, processes 7758-09-0, Potassium nitrite 7779-88-6, Zinc nitrate 10043-52-4, Calcium chloride, processes 10124-37-5, Calcium nitrate 10377-60-3, Magnesium nitrate

RL: DEV (Device component use); MOA (Modifier or additive use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (deliquescent particles; adhesive mixt. for fluorescent particle coating on cathode emitter tubes)

L52 ANSWER 32 OF 37 HCAPLUS COPYRIGHT 2002 ACS

AN 1994:691016 HCAPLUS

DN 121:291016

TI Cathode emitters and manufacturing thereof

IN Toyoda, Osamu; Betsui, Keiichi; Fukuda, Shinya; Nakatani, Tadashi

PA Fujitsu Ltd, Japan

SO Jpn. Kokai Tokkyo Koho, 11 pp. CODEN: JKXXAF

DT Patent

LA Japanese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΙ	JP 06020592	A2	19940128	JP 1992-334391	19921215
	JP 06016984	A2	19940125	JP 1992-334754	19921215
PRAI	JP 1992-113708		19920506		

Title manufg. involves (1) depositing a 1st material for forming a cathode-forming layer, (2) depositing a 2nd material for forming emitter tip neg.-feedback resistors, (3) depositing a 3rd material for forming emitter tip electron emitting edges, (4) forming masks over regions for formation of the emitter tips, (5) etching to give undercuts below the masks, and (6) subsequently forming an insulator film and a gate electrode film, wherein the 1st, 2nd, and 3rd materials are made of semiconductor materials. The manufg. gives simple prepn. of the emitter tip neg.-feedback resistors and low-resistance electron emission edges independently.

IT Cathodes

(emitter; manufg. for easy sep. formation of neg.-feedback

resistor and low-resistance electron emission edge for) Semiconductor materials (for formation of cathode and emitter tip neg.-feedback resistor and electron emitting edge in cathode emitter) Electric resistors (neg.-feedback; formation in manufg. cathode emitter) 7439-98-7, Molybdenum, processes 7440-25-7, Tantalum IT , processes RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (semiconductor; cathode electrode in cathode emitter) 7440-21-3, Silicon, processes IT RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses) (semiconductor; neg.-feedback resistor in cathode emitter) L52 ANSWER 33 OF 37 HCAPLUS COPYRIGHT 2002 ACS AΝ 1994:547007 HCAPLUS 121:147007 DN Field emission device and fabrication thereof TI IN Watanabe, Teruo; Ito, Shigeo; Ootsu, Kazuyoshi; Taniguchi, Masateru; Nishimura, Norio; Ochiai, Hisataka; Yamaguchi, Manabu PA Futaba Denshi Kogyo Kk, Japan Jpn. Kokai Tokkyo Koho, 4 pp. SO CODEN: JKXXAF DT Patent LΑ Japanese FAN.CNT 1 KIND DATE PATENT NO. APPLICATION NO. DATE ----A2 19931112 ΡI JP 05299011 JP 1992-99420 19920420 JP 2636630 B2 19970730 Title fabrication involves (1) forming a cathode conductor, an insulator, AΒ and a gate successively on an insulated substrate, followed by opening a hole to the insulator and the gate, (2) depositing a Si layer to form a resistor layer on the cathode exposed through the hole, (3) anodizing the Si layer to give a peeling layer followed by depositing **Mo** to give an **emitter** on the resistor layer, and (4) removing unnecessary portions of the deposition layer by removing it together with the peeling layer to form sep. resistors in every emitter formed on the device. The fabrication provides a no. of emitter/resistor units in the device simultaneously. Electric resistors IT (formation of, in emitter, in fabrication of field emission device) TT Electron emission (field, device, fabrication of, formation of resistor in) IT 7440-42-8, Boron, uses 7723-14-0, Phosphorus, uses RL: PEP (Physical, engineering or chemical process); PROC (Process) (dopant, in silicon resistor, for fabrication of field emission device) 7440-21-3, Silicon, uses IT RL: USES (Uses) (doped, resistor in emitter for field emission device, fabrication of) L52 ANSWER 34 OF 37 HCAPLUS COPYRIGHT 2002 ACS AN1994:546649 HCAPLUS DN 121:146649 TIMOS-controlling static induction thyristors

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Saito, Masayuki
IN
   Yazaki Corp, Japan
PA
    Jpn. Kokai Tokkyo Koho, 7 pp.
SO
    CODEN: JKXXAF
DT
    Patent
LA
   Japanese
FAN.CNT 1
                                       APPLICATION NO. DATE
                   KIND DATE
    PATENT NO.
    _____
                                         -----
    JP 06021437
                                        JP 1993-65604 19930324
                   A2 19940128
                          19920415
PRAI JP 1992-95259
    Title thyristor comprises a 1st cond.-type semiconductor substrate, 2nd
    cond.-type 1st and 2nd gate diffusion layers formed on the substrate, a
    1st cond.-type cathode diffusion layer formed between
    the gate diffusion layers, a 2nd cond.-type cathode-emitter
    -short diffusion layer, and a doped polysilicon layer formed via an
    insulator layer between the gate diffusion layer and
    cathode-emitter-short diffusion layer to give 1st and
    2nd cond.-type MOS transistors. The thyristor gives an improved
    switching characteristics without an increase of MOS
    capacitance.
    Thyristors
IT
       (MOS-controlling, static induction, improvement of switching
       characteristic without increase of MOS capacitance)
IT
    7440-21-3, Silicon, uses
    RL: USES (Uses)
       (doped, for gate electrode, in MOS-controlling thyristors)
L52 ANSWER 35 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    1993:593688 HCAPLUS
AN
DN
    119:193688
    Semiconductor devices having MOS anode short structure with
TI
    excellent turn-on-loss and turn-off-loss trade off
IN
    Shimizu, Naohiro
PΑ
    Toyo Electric Mfg Co Ltd, Japan
    Jpn. Kokai Tokkyo Koho, 11 pp.
SO
    CODEN: JKXXAF
DT
    Patent
LA
    Japanese
FAN.CNT 1
                   KIND DATE
                                        APPLICATION NO. DATE
    PATENT NO.
                          -----
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                                         -----
    JP 05095112 A2 19930416
JP 3160330 B2 20010425
PΙ
                          19930416
                                         JP 1991-280646 19911001
    In a bipolar semiconductor device comprising a laminate of a
AΒ
    cathode n-emitter layer, a p-base layer or
    p-gate layer, a high-resistance semiconductor layer, an n-buffer layer,
    and an anode p-emitter layer, the anode p-emitter
    layer is etched, an MOS anode short layer which works as a short
    layer is arranged in such a way that it contacts with the anode p-
    emitter layer facing the etched groove but not contacting the
    n-buffer layer, the space between the etched groove and part of the
    MOS anode short layer which works as a n+ short layer is coated
    with a SiO2 film and then coated with a metal. The trade off of
    turn-on-loss and turn-off-loss is improved.
IT
    Semiconductor devices
       (MOS anode short arrangement, with excellent trade off of
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turn-on-loss and turn-off-loss)

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L52 ANSWER 36 OF 37 HCAPLUS COPYRIGHT 2002 ACS
    1978:82762 HCAPLUS
AN
DN
    88:82762
ΤI
    Gold-diffused thyristor
TN
    Moyson, Joseph
    General Electric Co., USA
PΑ
    U.S., 5 pp.
SO
    CODEN: USXXAM
DT
    Patent
LA
    English
FAN.CNT 1
                    KIND DATE
                                         APPLICATION NO. DATE
    PATENT NO.
     _____
                                          _____
PΙ
    US 4066484
                     Α
                           19780103
                                          US 1975-616404 19750924
                           19741024
PRAI US 1974-517523
     Au-doped fast turnoff thyristor is manufd. conventionally
AB
     through the diffusion of the cathode emitter
     layer. Following diffusion of the cathode
     emitter layer, the diffusion mask is left in place and
     an auxiliary dopant that stimulates carrier recombination is diffused into
     the body. Use of the cathode emitter mask ensures that the
     auxiliary dopant diffuses only into the cathode emitter and the
     area there around, where it is most effective in enhancing fast turnoff
     characteristics, but does not contribute substantially to excess leakage
     current. Moreover, utilization of a preexisting mask minimizes the cost
     involved in the Au doping process. Both Au-diffused
     fast turnoff triacs and semiconductor controlled rectifiers can be made by
     substantially the same method.
IT
    Thyristors
        (silicon, gold-diffused)
     7440-57-5, uses and miscellaneous
IT
     RL: PEP (Physical, engineering or chemical process); PROC (Process)
        (doping with, of silicon thyristors for rapid turnoff)
     7440-21-3, uses and miscellaneous
TT
    RL: USES (Uses)
        (thyristors, gold-diffused)
L52
    ANSWER 37 OF 37 HCAPLUS COPYRIGHT 2002 ACS
     1972:65270 HCAPLUS
AN
DN
     76:65270
     Electron conduction, electron emission, and electroluminescence of MIM
ΤI
     [metal-insulator-metal] sandwich structures with aluminum(III) oxide
     insulating layers
ΑU
     Dittmer, G.
CS
     Philips Forschungslab. Aachen G.m.b.H., Aachen, Ger.
     Thin Solid Films (1972), 9(2), 141-72
SO
     CODEN: THSFAP
DT
     Journal
LA
     English
     The elec. properties of Al-Al203-Au sandwiches with different
AB
     Al203 structures were investigated. Sandwich cathodes with
     Al203 layers .ltoreq.100 .ANG. thick, or with thicker thermally
     oxidized or anodically grown dense Al203 layers, give monotonic
     current-voltage characteristics. The lifetime of such cathodes is limited
     to a few min by uniform Au ion diffusion at higher emission
     yields. Sandwich cathodes having a multilayered Al2O3 structure with at
     least one heavily doped porous Al2O3' layer exhibit current-voltage curves
     with voltage-controlled neg. resistance. At >5 V, the sandwich and the
     emission currents may be attributed to Fowler-Nordheim tunnelling.
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electron-emitting regions of the sandwich structure show faint electroluminescence with peaks of emitted photons at 2.3 and 4.1 eV. From a consideration of different electron conduction mechanisms, related to obsd. elec. cond., electron emission, and electroluminescence, a band structure was derived for amorphous Al2O3' decorated with Au pptns. The investigations led to the construction of a sandwich cathode with extended lifetime. The main features of this cold emitter are a special multilayer structure of the Al2O3 insulator and a discontinuous upper Au electrode.

IT Cathodes

(aluminum-aluminum oxide-gold sandwich structures as, lifetime of)

IT Luminescence

(electro-, of aluminum-aluminum oxide-gold sandwich structures)

IT Electron emission

(from aluminum-aluminum oxide-gold sandwich structures)

IT Energy level, band structure

(of aluminum oxide, in sandwich structures with aluminum and gold)

IT Electric conductivity and conduction

Electric current-potential relationship

(of aluminum-aluminum oxide-gold sandwich structures)

IT 1344-28-1, properties

RL: PRP (Properties)

(elec. insulation by and elec. properties of, in sandwich structures with aluminum and $\ensuremath{\mathbf{gold}}\xspace)$